

KW treatment: Neisseria infection; meningitis; septicaemia; gonorrhea.
 XX OS Neisseria meningitidis.

XX PN W09924578-A2.

XX PD 20-MAY-1999.

XX PF 09-OCT-1998; 98W0-IB01665.

XX PR 01-SEP-1998; 98GB-0019016.

XX PR 06-NOV-1997; 97GB-0023516.

XX PR 14-NOV-1997; 97GB-0024190.

XX PR 18-NOV-1997; 97GB-0024386.

XX PR 27-NOV-1997; 97GB-0025158.

XX PR 10-DEC-1997; 97GB-0026147.

XX PR 14-JAN-1998; 98GB-0000759.

XX PA (CHIR-) CHIRON SPA.

XX PI Grandi G, Maignani V, Pizza M, Rappelli R, Scarlato V;

XX DR WPI: 1999-327407/27.

XX DR N-PSDB; AA212027.

XX PT Proteins from Neisseria meningitidis and N. gonorrhoeae useful for

XX PT diagnosis, treatment and prevention of infection

XX PS Claim 4; Page 123; 524pp; English.

XX SO Sequence 447 AA;

alignment_scores:

Quality: 2208.00

Ratio: 5.088

Percent Similarity: 97.092

Align seg 1/1 to: AAY38562 from: 1 to: 447

Length: 447
Gaps: 0

Percent Identity: 97.092

alignment_block:

US-09-303-518d-127 x AAY38562

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17 uGlnValIleuysArgIleuysIleuysIleuysIleuysIleuysG 34
101 AAGATATGCGCGATGCGCGCGCTMGATGAAGTCAAGAGGCGATGCC 150
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34 InGIuylrAlaGlyMetArgPro**MetIleuysValIleuysIleuys 50
151 GTCAAAAAGGCCAAGTGTGTTGAAGCAAAAGNATCCGGGGTGTGT 200
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51 ValIleuysIleuysIleuysIleuysIleuysIleuysIleuys 67
201 GTTTACCGCGCGCGTTCAGCAAAATCGCGCATCCATCGCGCGAAA 250
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351 ANTNGNNGCAATCTGATCCATCCGGTGTGTGACTGCGCTGTANCC 400
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401 GTCCGTTACGCAAAATCCCTGCGCGTGCAGAGCGCGCGTCCGATTC 450
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134 rgProPheSerIleProAlaValAspIleuArgIleProPheAlaIle 150
451 GTCAATCGATGACACCAATCCGCTNCGCGGACGACCCCTGTGTGAT 500
|||||
151 ValAsnAlaMetAspThrAsnProLeuAlaIleAspProValValIle 167
501 CAAGAAGCGCGANGATTTTCAGACGANGTNTGTGTATGAGCCGTT 550
|||||
167 elysGluAla*****AspPheArgArg*****LeuValIleuSerArg 184
551 TGACCGAGCGTAAATCCATGTGTGTAGGCGACGCTGCGCAGCGTCC 600
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601 TCTGAATGCTGCGCAACATCCATGCAACATGATTCGGCGCGCGATCC 650
|||||
201 SerIleuAsnAlaIleAsnIleGluThrHisGluPheGlyProHisPr 217
651 GCGCGGTTAGTGGCGACGACATTCATTTCATGAGCGCGTGGTGC 700
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951 GGAAGCGCTACCAATCAGATTCCTGTTATCGAAGAGCGCGCGAAG 1000
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1251 AGACCTGCTTTGTCAGCTTCGTGCGCCGGCAAAATACGAATAGCC 1300
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seq_documentation_block:
ID AAV38561 standard; Protein; 447 AA.
AC AAV38561;
XX
XX 08-OCT-1999 (first entry)
DE Neisseria meningitidis antigen encoded by ORF22.
XX
XX Neisseria meningitidis; Neisseria gonorrhoeae; antigen; vaccine;
KM treatment; Neisseria infection; meningitis; septicemia; gonorrhea.
XX
OS Neisseria meningitidis.
XX
XX WO924578-A2.
XX
XX 20-MAY-1999.
XX
XX 09-OCT-1998; 98WO-IB01665.
XX
XX 01-SEP-1998; 98GB-0019016.
XX
XX 06-NOV-1997; 97GB-0023516.
XX
XX 14-NOV-1997; 97GB-0024190.
XX
XX 18-NOV-1997; 97GB-0024386.
XX
XX 27-NOV-1997; 97GB-0025158.
XX
XX 10-DEC-1997; 97GB-0026147.
XX
XX 14-JAN-1998; 98GB-0000759.
XX
XX (CHIR-) CHIRON SPA.
XX
XX Grandi G, Masignani V, Pizza M, Rappuoli R, Scarlato V;
XX WPI: 1999-327407/27.
XX
XX N-PSDB; AA212026.
XX
XX Proteins from Neisseria meningitidis and N. gonorrhoeae useful for
XX diagnosis, treatment and prevention of infection
XX
XX Claim 4; Page 123; 524pp; English.
XX
XX Amino acid sequences AAV38499-138944 represent Neisseria meningitidis
XX and N. gonorrhoeae antigenic proteins. They are encoded by open
XX reading frames (ORFs) AA211972-212358. The antigenic proteins,
XX their fragments, their nucleic acids and antibodies are used for
XX diagnosis, prevention (as vaccines) or treatment of Neisseria
XX infections, such as meningitis, septicemia and gonorrhea. Both
XX organisms are closely related. Fragments of the nucleic acids
XX are useful as hybridisation probes and antisense reagents.
XX
XX Sequence 447 AA:

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alignment_scores:
  Quality: 2177.00      Length: 447
  Ratio: 4.982          Gaps: 0
  Percent Similarity: 97.763  Percent Identity: 94.855

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17 uGlnAlaValTyrAspGlyProAlaIleThrGluValAlaLeuLeuGlyG 34
101 AACAATATGCGCGGTATGCGCCCTTGATGAAATGCAAGAGAGCGATGC 150
34 IuGluTyrAlaGlyMetArgProSerMetLysValLysGluGlyAspAla 50
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51 ValLysLysGlyGlnValLeuPheGluAspLysLysAsnProGlyValVal 67
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67 IPhetIrrAlaProAlaSerGlyLysIleAlaIleAlaIleHisArgGlyL 84
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134 rGProPheSerLysIleProAlaValAlaSprAlaGluProPheAlaIleP 150
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651 GCGCGGTTTGAAGGACGACGACATTCATTCAATGACCGGCTGGTCAA 700
217 oAlaGlyLeuSerGlyThrHisIleHisPheIleGluProValGlyAla 234
701 ACAAAACCGTTGGACCATCAATTAACAGATGAATGATCAATGACGAG 750
234 snLysThrValIleThrHisAsnTyrGlnAspAlaIleThrIleGlyArg 250
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434 rLeuLeuArgGlyValLeuGluThrIleGluLysGluGly 447
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seq_documentation block:
ID AA198564 standard; Protein; 447 AA.
XX
AC AA198564;
XX
DT 08-OCT-1999 (first entry)
XX
DE Neisseria gonorrhoeae antigen encoded by ORF22.
XX
KW Neisseria meningitidis; Neisseria gonorrhoeae; antigen; vaccine;
treatment; Neisseria infection; meningitis; septicemia; gonorrhea.
OS Neisseria gonorrhoeae.
XX
PN WO9924578-A2.
XX
PD 20-MAY-1999.
XX
PF 09-OCT-1998; 98MO-IB01665.
XX
PR 01-SEP-1998; 98GB-0019016.
PR 06-NOV-1997; 97GB-0023516.
PR 14-NOV-1997; 97GB-0024190.
PR 18-NOV-1997; 97GB-0024386.
PR 27-NOV-1997; 97GB-0025158.
PR 10-DEC-1997; 97GB-0026147.
PR 14-JAN-1998; 98GB-0000759.
XX
XX (CHIR-) CHIRON SPA.
XX
PA Grandi G, Malignani V, Piazza M, Rappuoli R, Scarlato V;

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XX
DR WPI: 1999-327407/27.
N-PSDB; AA12028.
XX
PT Proteins from Neisseria meningitidis and N. gonorrhoeae useful for
diagnosis, treatment and prevention of infection
XX
PS Claim 4; Page 125; 524pp; English.
XX
CC Amino acid sequences AA198499-Y38944 represent Neisseria meningitidis
and N. gonorrhoeae antigenic proteins. They are encoded by open
reading frames (ORFs) AA11972-212358. The antigenic proteins,
their fragments, their nucleic acids and antibodies are used for
diagnosis, prevention (as vaccines) or treatment of Neisseria
infections, such as meningitis, septicemia and gonorrhea. Both
organisms are closely related. Fragments of the nucleic acids
are useful as hybridisation probes and antisense reagents.
XX
SQ Sequence 447 AA;

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Quality: 2148.00 Length: 447
Ratio: 4.927 Gaps: 0
Percent Similarity: 97.539 Percent Identity: 93.289

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alignment_block:
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Align seg 1/1 to: AA198564 from: 1 to: 447

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51 GCAAGTCATTTATGACGGCGCGTCACTTACCGAAGTCCGCTTGGCG 100
|||||
17 uGlnValIleTyrAspGlyProAlaIleThrGluValAlaLeuLeuGlyG 34
|||||
101 AACAATATGCGGGTATGCGCGCGCGCGCGCGCGCGCGCGCGCGCG 150
|||||
34 IuGluTyrValGlyMetArgProSerMetLysIleLysGluGlyAla 50
|||||
151 GTCAAAAAGGCCCAAGTCTGTTGAAGACAAAAGNATCCGGCGGTGT 200
|||||
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201 GTTTACCGCGCGCGTTCAGGCAAAATCCCGCCATCCATCCGCGGAA 250
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67 IPhetThrAlaProAlaSerGlyLysIleAlaIleHisArgGlyGlu 84
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117 sValArgArgAsnLeuIleGlnSerGlyLeuThrPThrAlaLeuArgThrA 134
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451 GTCAATGCGATGACACCAATCCGCTNGCGGACGCGCGTGGTGTGAT 500
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201 SerGluAsnAlaAlaAsnIleGluThrHisGluPheGlyGlyProHisI 217
651 GCGCGGTTGAGTGGCAGCGACATTTATTCATTTGACCGCGTGGTCAA 700
217 oAlaGlyLeuSerGlyThrHisIleHisPheIleGluProValGlyAla 234
701 ACAAAACCGTTTGGACCATCAATTATCAATGATTAATTCCTGGAGCT 750
234 snLysThrValTrpThrIleAsnTyrGluAspValIleAlaIleGlyArg 250
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251 LeuPheValThrGlyArgLeuAsnThrGluArgValAlaIleGluGly 267
801 TTCTCAAGTCAACAAACCGCGCTGTGGTACCGTTTGGTGGTGGAAAG 850
267 yLeuGluValAsnLysProArgLeuAsnArgThrValLeuGlyAlaLys 284
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284 aISerGluLeuThrAlaGlyGluLeuValAspAlaAspAsnArgValIle 300
901 TCCGGTGTGATTTGACGCGCGGATTCACACAGCGCGGACGATTAATT 950
301 SerGlySerValLeuAsnGlyAlaIleAlaGlnGlyAlaHisAspTyrLe 317
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1051 AGACCCCTCGGCGCATTTCTGAAAAACAACCTTCAGTTCACGACAGC 1100
351 ThrThrLeuGlyHisPheLeuLysAsnLysLeuPheLysPheThrTrAl 367
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367 aValAsnGlyGlyAspArgAlaMetValProIleGlyThrTyrGluArg 384
1151 TAATGCCGCTAGACATCTGCTGCTGCTGCTTGGCGGATTAATGCTC 1200
384 alMetProLeuAspIleLeuProThrLeuLeuAsnArgAspLeuIleVal 400
1201 GCGCATCCGACAGCGCGCAAGCATTTGGTGTGGAATGGACGAAGA 1250
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ID AAV38563 standard; Protein: 322 AA.
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XX AAV38563;
XX

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DT 08-OCT-1999 (first entry)
XX
DE Neisseria gonorrhoeae antigen encoded by a partial ORF22.
XX
KW Neisseria meningitidis; Neisseria gonorrhoeae; antigen; vaccine;
XX treatment; Neisseria infection; meningitis; septicemia; gonorrhea.
XX
OS Neisseria gonorrhoeae.
XX
PN W0924578-A2.
XX
PD 20-MAY-1999.
XX
PF 09-OCT-1998; 98WO-IB01665.
XX
PR 01-SEP-1998; 98GB-0019016.
XX 06-NOV-1997; 97GB-0023516.
XX 14-NOV-1997; 97GB-0024180.
XX 18-NOV-1997; 97GB-0024386.
XX 27-NOV-1997; 97GB-0025158.
XX 10-DEC-1997; 97GB-0026147.
XX 14-JAN-1998; 98GB-0000759.
XX
PA (CHIR-) CHIRON SPA.
XX
PI Grandi G, Masignani V, Pizza M, Rappuoli R, Scarlato V;
XX WPI; 1999-327407/27.
XX
PT Proteins from Neisseria meningitidis and N. gonorrhoeae useful for
XX diagnosis, treatment and prevention of infection
XX
PS Claim 4; Page 124-125; 524pp; English.
XX
CC Amino acid sequences AAV38499-Y38944 represent Neisseria meningitidis
XX and N. gonorrhoeae antigenic proteins. They are encoded by open
XX reading frames (ORFs) Aa11972-212358. The antigenic proteins,
XX their fragments, their nucleic acids and antibodies are used for
XX diagnosis, prevention (as vaccines) or treatment of Neisseria
XX infections, such as meningitis, septicemia and gonorrhea. Both
XX organisms are closely related. Fragments of the nucleic acids
XX are useful as hybridisation probes and antisense reagents.
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SQ Sequence 322 AA:
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Percent Similarity: 97.205 Percent Identity: 91.304
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17 uGluValIleTyrAspGlyProAlaIleThrGluValAlaLeuLeuGly 34
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301 GAATGCAACGCTACGCGCCGCAAGCGTTGGCAAACTTAAGCGCGANGA 350
101 GluPheGluArgTyrValProGluAlaLeuAlaLysLeuSerSerGlyL 117
351 ANNNNGNCAATCATCTCAATCCGCTTGTGAGCGCGCTGCGCTANCC 400
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134 rGProPheSerLysIleProAlaValAspAlaGluProPheAlaIlePhe 150
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501 CAAAGAGCCGCGANGATTTCAGACGANGTGTCTGATTAGCCGCTT 550
167 elysGluAlaIleAlaAspPheLysArgGlyLeuLeuValLeuSerArgL 184
551 TGACCGAGCGTAAATCATCTGTGTGAAGCAGCTGCGCAGACGTCCG 600
184 eutThrGluArgLysIleHisValCysLysAlaIleGlyAlaAspValPro 200
601 TCTGAATGCTGCCACATCGAAACACATGATTCGGCGCGCCGATTC 650
201 SerGluAsnAlaIleAsnIleGluThrHisGluPheGlyGlyProHisr 217
651 GCGCGTTTGAATGCGCAGCAGCATTCATTTCATTGACCGCGTGTGCA 700
217 calAGlyLeuSerGlyThrHisIleHisPheIleGluProValGlyAla 234
701 ACAAAACGTTTGACCATCATATATCAAGATTAATTCGACGCGACGT 750
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267 yLeuGlnValAsnLysProArgLeuLeuArgThrValLeuGlyAlaLysV 284
851 TATTCGAATTAAGTGGCGGCGAATTTGGTGAAGCAGCAGCAGCGAT 900
284 alSerGlnLeuThrAlaGlyLeuValAspAlaAspAsnArgValIle 300
901 TCCGTTGCTGATTTGAACGCGCGATTACACAGCGCGCAGCATTTAT 950
301 SerGlySerValLeuAsnGlyAlaIleAlaGlnGlyAlaHisAspTyrLe 317
951 GGGAGCGTACCAAT 966
317 uGlyArgTyrHisAsn 322
seq_name: /SISL/gcgdata/geneseq/geneseq-emb1/AA1999.DAT:AA1999
seq_documentation_block:
ID AA1999 standard; Protein: 158 AA.
XX
AC AA1999:
XX
DT 08-OCT-1999 (first entry)
XX
DE Neisseria meningitidis antigen encoded by a partial ORF22.

```

```

XX
KM Neisseria meningitidis; Neisseria gonorrhoeae; antigen; vaccine;
KN treatment; Neisseria infection; meningitis; septicemia; gonorrhea.
XX
OS Neisseria meningitidis.
XX
PN MO924578-A2.
XX
PD 20-MAY-1999.
XX
PF 09-OCT-1998; 98MO-IB01665.
XX
PR 01-SEP-1998; 98GB-0019016.
PR 06-NOV-1997; 97GB-0023516.
PR 14-NOV-1997; 97GB-0024190.
PR 18-NOV-1997; 97GB-0024386.
PR 27-NOV-1997; 97GB-0025158.
PR 10-DEC-1997; 97GB-0026147.
PR 14-JAN-1998; 98GB-0000759.
XX
PA (CHIR-) CHIRON SPA.
XX
PI Grandi G, Masignani V, Pizza M, Rappuoli R, Scarlato V;
XX WPL 1999-327407/27.
XX DR N-PSDB; AA212025.
XX
PT Proteins from Neisseria meningitidis and N. gonorrhoeae useful for
XX diagnosis, treatment and prevention of infection
XX
PS Claim 4; Page 122; 524pp; English.
XX
ES Amino acid sequences AA1999-138944 represent Neisseria meningitidis
XX CC and N. gonorrhoeae antigenic proteins. They are encoded by open
XX CC reading frames (ORFs) AA211972-212358. The antigenic proteins,
XX CC their fragments, their nucleic acids and antibodies are used for
XX CC diagnosis, prevention (as vaccines) or treatment of Neisseria
XX CC infections, such as meningitis, septicemia and gonorrhea. Both
XX CC organisms are closely related. Fragments of the nucleic acids
XX CC are useful as hybridisation probes and antisense reagents.
XX
SQ Sequence 158 AA:
XX
alignment_scores:
Quality: 739.00 Length: 158
Ratio: 4.862 Gaps: 0
Percent Similarity: 96.203 Percent Identity: 92.405
alignment_block:
US-09-303-518D-127 x AA1999
Align seg 1/1 to: AA1999 from: 1 to: 158
1 ATGATTAATCAAAAGGCTTAACCTGCGCGGCGGAGCGGAGCGGA 50
1 MetIleLysIleLysGlyLeuAsnLeuProIleAlaIleArgProG 17
51 GCAAGTCAATTAAGCGCGCGCGCTTACCGAAGTCCGCTTCTTGGG 100
17 uGlnAlaValTyrAspGlyProAlaIleThrGluValAlaLeuLeuGly 34
101 AAGATTAATGCGCGGTATGCGCGCGCGCGCGCGCGCGCGCGATG 150
34 uGluTyrAlaGlyMetArgProSerMetLysValLysGluGlyAspAla 50
151 GTCAAAAGGCGCAAGTCTGTTTGAAGACAAAGATCCGCGCGTGGT 200
51 ValLysLysGlyGlnValLeuPheGluAspLysAsnProGlyValVa 67
201 GTTAACGCGCGCGCGTTCAGGCAAAATCGCGCGCATCGCGGCGAA 250
67 lPheThrAlaProAlaSerGlyLysIleAlaIleHisArgGlyIuL 84

```

```

251 AGCGGCTACTTCAGTCGCTGCTGATTGCCGTTGAAGCAGCAGCAAAATC 300
|||||  |||||||  |||||||  |||||||  |||||||  |||||||  |||||||
84 ysarGValLeuInSerValValIleAlaValGlu**AsnAspGluIle 100
301 GAGTTGCAAGCCTACGCGCCGCAAGCGTTGGCAAACTTAAGCGCGGANGA 350
|||||  |||||||  |||||||  |||||||  |||||||  |||||||  |||||||
101 GIuphegiuArgTyraIaProGIuAlaLeuAlaAsnLeuSerGIyGIuGI 117
351 ANTNGNNGCAATCGATCCAAATCCGTTTGTGACTGCCGCTGGGTANCC 400
|||||  |||||||  |||||||  |||||||  |||||||  |||||||  |||||||
117 uValArgArGAsnLeuIleGIuInSerGIyLeuTrpThrAlaLeuArgTrp 134
401 GTCCGTTGACAAAATCCCTGCGCTGATGCCGAGCGCGCTTGCATCTTC 450
|||||  |||||||  |||||||  |||||||  |||||||  |||||||  |||||||
134 rGProPheSerLysIleProAlaValAlaAspAlaGIuProPheAlaIlePhe 150
451 GTCAATGCGATGACACCAATCCG 474
|||||  |||||||  |||||||  |||||||  |||||||  |||||||  |||||||
151 ValAsnAlaMetAspThrAsnPro 158

```

seq_name: /SIDSL/gcgdata/geneseq/geneseq-emb1/AA1999.DAT:AAV34439

seq_documentation_block:

ID AAV34439 standard; Protein; 451 AA.

XX AAV34439;

XX 25-AUG-1999 (first entry)

XX Porphyromonas gingivalis protein PGI.

XX Porphyromonas gingivalis; PG; periodontal disease; gingivitis;

KW vaccine; antigenic.

OS Porphyromonas gingivalis.

XX WO929870-A1.

XX 17-JUN-1999.

XX 10-DEC-1998; 98WO-AU01023.

XX 04-AUG-1998; 98AU-0005028.

XX 10-DEC-1997; 97AU-0000839.

XX 31-DEC-1997; 97AU-0001182.

XX 30-JAN-1998; 98AU-0001546.

XX 10-MAR-1998; 98AU-0002264.

XX 09-APR-1998; 98AU-0002911.

XX 23-APR-1998; 98AU-0003128.

XX 05-MAY-1998; 98AU-0003338.

XX 22-MAY-1998; 98AU-0003654.

XX 29-JUL-1998; 98AU-0004917.

XX (CSLC-) CSL LTD.

XX Agius CT, Barr IG, Hocking DM, Margetis MB, Patterson MA;

XX PI Ross BC, Rothel LJ, Webb EA;

XX WPI; 1999-385613/32.

XX N-PSDB; AAV31657.

XX Antigenic Porphyromonas gingivalis peptides for preventing

CC be used to detect Porphyromonas gingivalis in standard hybridisation

CC assays. Porphyromonas gingivalis is involved in periodontal disease

XX especially gingivitis.

XX SQ Sequence 451 AA;

alignment_scores: Quality: 636.00 Length: 452
Ratio: 2.208 Gaps: 7
Percent Similarity: 63.717 Percent Identity: 34.071

alignment_block:

US-09-303-518d-127 x AAV34439 ..

Align seg 1/1 to: AAV34439 from: 1 to: 451

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1 ATGATTAATAAATCAAAAAAGCTTAAACCTGCCCATCGCGGAGACCG.. 48
:::|||||  |||||||  |||  ::  ||::|||
4 ValIleLysThrLysLysGIyLeuAlaLeuAlaAsnLeuLysGIyLysPro 20
49 .GAGCAAGTCATTTATGACGGCGCCGTCATTTACCGAAGTCGGCTCTTG 97
:::|||||  ::  ||::|||
20 uProGIuMetLeuAlaGIuProAlaGIuInSerProThrTyraIaValAla 37
98 GCGAAGATATGCCGATGCGCCGCTGATGCCGCTTNGATGAAAGTCAAGAGCGAT 147
:::|||||  ||::|||  |||  ::  ::::  |||||
37 roAspAspPheGIuGIyValIleProLysValIleAlaArgProGIyAsp 53
148 GCCGTCAAAAAAGCCCAAGTCGTGTTGAAGCAAAAAAGNATCCGGCGCT 197
||||:  ||::|||  ||::|||  |||  ::  |||
54 LysValArgAlaGIySerAlaLeuMetHisLysAlaTyLysProGIuMet 70
198 GGTGTTACCGCGCCGCTTTCAGGCAAAATCGCGCCATCCATCGCGCG 247
:  ||::|||  ||::|||  ||::|||  ||::|||  ||::|||
70 LysPheThrSerProValSerGIyGIuValIleAlaValAsnArgGIyA 87
248 AAAAGCGGCTACTTCAGTCGCTGATGCCGCTTGAAGCAAGAGCGAA 297
|||||  ::  ||::|||  ::  ||::|||
87 IalysArgLysValIleuSerIleGIuValLysProAspGIyLeuAsnGIu 103
298 ATCGAG...TTGAAACGCTACGCGCCGAGCGCTTGCAAACTTAAGCG 344
|||  |||  ||::|||
104 TyrGIuSerPheProValGIyAspProSerAla.....LeuSerAl 117
345 CGANGAANTNGNNGCAATCTGATCCATCCGCTTGAGACTGGCGCTCG 394
:  ::::  ||::|||  ||::|||  ||::|||
117 aGIuGIuIleLysGIuLeuLeuLeuSerGIyMetTrpGIyPheIleu 134
395 GTANCCGTCGCTTCAGCAAAATCCCTGCGCTGATGCCGAGCGCTGCGC 444
::  ||::|||  ::  ::  |||
134 LysGIuArgProTyraPheIleValAlaIleThrProAspIleAlaProAla 150
445 ATCTTCGTCATGCGATGACACCAATCCGCTNGCGGAGACCGCTGTG 494
||||:  ||::|||  |||  |||||  |||
151 IleTyrIleThrAlaAsnPheThrAlaProLeuAlaProAspPheAsp 167
495 TGGATCAAGAAGCGCGANGATTTGAGACGANGANTNGCTGTGTTGA 544
:::|||||  ::  ||::|||  ::  ||::|||
167 eIleValArgGIyGIuGIuArgAlaLeuGIuInThrAlaIleAspAlaLeu 184
545 GCCGTTTGACCGAGCGTAAATCCATGCTGTAAGCAGACGCGCGAGAC 594
:::|||||  ||::|||  ||::|||  ||::|||
184 IalysLeuThrTrpGIyLysValTyValGIyLeuLysProGIySerSer 200
595 GTGCCGCTGAATAATGCTGCCAATCGAACAACATGAATTCGGCGCGCC 644
::  ||::|||  |||
201 LeuGIyLeuHisAsnAlaGIuIleValGIuValHis.....GIyPr 214
645 GCATCCGCGCGGCTTGAGTGCGACGCAATTCATTTCATTGAGCGCGTG 694
|||||  ||::|||  ||::|||  ||::|||
214 OHISProAlaGIyAsnValGIyValLeuIleAsnHisThrLysProIleA 231

```


CC Neisserial bacteria (e.g. meningitis and septicaemia), to detect the
 CC presence of Neisseria bacteria, or to raise antibodies. They may also
 CC be used to screen for agonists or antagonists, which may themselves
 CC have use as antibacterial agents. The polynucleotides of the invention
 CC may also be used in gene therapy protocols.

XX Sequence 120 AA;

alignment_scores:

Quality: 563.00 Length: 120
 Ratio: 4.896 Gaps: 0
 Percent Similarity: 95.833 Percent Identity: 94.167

alignment_block:

US-09-303-518D-127/rev x AAY75273

Align seg 1/1 to: AAY75273 from: 1 to: 120

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674 ATGTGCGTGCACCTCAACCGCGCGGATCGGCGCCGCGCATTCATGTGT 625
|||||
1 MetCysValProLeuYsProAlaGlyCysGlyProProAsnSerCysVa 17
624 TTTCGATGTGGCAGCATTTTCAGACGGCAGCTCTGCGCCAGCTGCTTAC 575
|||||
17 lserMetLeuAlaAlaPheSerAspGlyThrSerAlaProAlaAlaLeuH 24
574 ACACATGATTTTACGCTCGGTCAACCGGCTCAATTCACGACNACNTGCT 525
|||||
34 lserThrPileuAlaArgSerValYsArgLeuAsnThrSerYsProArg 50
524 CTGAATATCNCNCGCGCTTTTGTATCACACACGAGGCTGCGCGNAG 475
|||||
51 LeuYsSerSerAlaAlaSerLeuIleThrThrGlySerAlaAla 67
474 CGGATTTGGTGCATTCGATTCAGCAAGATGGCGAAGGCTTGCGATCGA 425
|||||
67 rGlyLeuValSerIleAlaLeuThrIlysmetAlaAsnGlySerAlaSerT 84
424 CGGACGAGGATTTTGTGAACGAGCGGNTACGACGCGAGTCCACAAACCG 375
|||||
84 hrAlaGlyIleLeuLeuAsnGlyArgValArgSerAlaValAlaHisYsPro 100
374 GATTGGATCAGATTGCGNCCNANTTCNTCCGCGCTTAAGTTGCCAAGCG 325
|||||
101 AspTrpIleArgLeuArgArgThrSerSerProLeuYsPheAlaAsnAl 117
324 TTTCGGCGCGG 315
|||||
117 aserGlyAla 120

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seq_name: /SID81/gcgdata/geneseq/geneseqp-emb1/AA2000.DAT:AAY75272

seq_documentation_block:

ID AAY75272 standard; Protein: 120 AA.

XX AAY75272;

DT 21-MAR-2000 (first entry)

DE Neisseria meningitidis ORF 628 protein sequence SEQ ID NO:2018.

XX Neisseria meningitidis; Neisseria gonorrhoeae; antigen; vaccine;

KW antigenic; diagnosis; immunogenic; infection; meningitis; septicaemia;

XX antibacterial; gene therapy.

XX Neisseria meningitidis.

XX WO957280-A2.

XX 11-NOV-1999.

XX 30-APR-1999; 99WO-US09346.

XX 01-MAY-1998; 98US-0083758.
 PR 31-JUL-1998; 98US-0094869.
 PR 02-SEP-1998; 98US-0098894.
 PR 02-SEP-1998; 98US-0099062.
 PR 09-OCT-1998; 98US-0103749.
 PR 09-OCT-1998; 98US-0103794.
 PR 09-OCT-1998; 98US-0103796.
 PR 25-FEB-1999; 99US-0121528.

XX (CHIR) CHIRON CORP.
 PA (GENO-) INST GENOMIC RES.

PI Fraser C, Galeotti C, Grandi G, Hickey E, Masignani V, Mora M;
 PI Petersen J, Pizsa M, Rappuoli R, Ratti G, Scalatò E, Scarselli M;
 PI Tettelin H, Venter JC;

DR WPI; 2000-062150/05.
 DR N-PSDB; AA254034.

PT Novel Neisserial polypeptides predicted to be useful antigens for
 PT vaccines and diagnostics
 PS Claim 2; Page 1003; 1453pp; English.

XX AA253015 to AA254536, AA254577 to AA254615, and AAY74253 to AAY75941
 CC represent novel Neisseria meningitis and N. gonorrhoeae polynucleotides
 CC and polypeptides. AA254537 to AA254576 and AA254616 to AA255473 represent
 CC PCR primers used in the exemplification of the present invention. The
 CC polypeptides, the polynucleotides, antibodies and compositions of
 CC the invention can be used as vaccines, as diagnostic reagents, and as
 CC immunogenic compositions. The polypeptides can be used in the
 CC manufacture of medicaments for treating or preventing infection due to
 CC Neisserial bacteria (e.g. meningitis and septicaemia), to detect the
 CC presence of Neisseria bacteria, or to raise antibodies. They may also
 CC be used to screen for agonists or antagonists, which may themselves
 CC have use as antibacterial agents. The polynucleotides of the invention
 CC may also be used in gene therapy protocols.

XX Sequence 120 AA;

alignment_scores:
 Quality: 536.00 Length: 120
 Ratio: 4.702 Gaps: 0
 Percent Similarity: 95.000 Percent Identity: 90.000

alignment_block:

US-09-303-518D-127/rev x AAY75272

Align seg 1/1 to: AAY75272 from: 1 to: 120

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674 ATGTGCGTGCACCTCAACCGCGGATCGGCGCCGCGCATTCATGTGT 625
|||||
1 MetCysValProLeuYsProAlaGlyCysGlyProProAsnSerCysVa 17
624 TTTCGATGTGGCAGCATTTTCAGACGGCAGCTCTGCGCCAGCTGCTTAC 575
|||||
17 lserMetLeuAlaAlaPheSerAspGlyThrSerAlaProAlaAlaLeu 34
574 ACACATGATTTTACGCTCGGTCAACCGGCTCAATTCACGACNACNTGCT 525
|||||
34 lserThrPileuAlaArgSerValYsArgLeuAsnThrSerYsProArg 50
524 CTGAATATCNCNCGCGCTTTTGTATCACACGAGGCTGCGCGNAG 475
|||||
51 LeuYsSerSerAlaAlaSerLeuIleMetThrValGlySerAlaAla 67
474 CGGATTTGGTGCATTCGATTCAGCAAGATGGCGAAGGCTTGCGATCGA 425
|||||
67 rGlyLeuValSerIleAlaLeuThrIlysmetAlaAsnGlySerAlaSerT 84
424 CGGACGAGGATTTTGTGAACGAGCGGNTACGACGCGAGTCCACAAACCG 375

```

```

|||||
84 hrAlaGlyIleLeuLeuAsnGlyArgValArgSerAlaValHisLysPro 100
374 GATTGGATTCAGATTCGNNCANNANTTCNTGCCGCTTACGTTGGCAACGC 325
|||||
101 AspIrrPileArgLeuArgArgThrSerSerProLeuLysPheAlaSerAl 117
324 TTCGGGGCGCG 315
|||||
117 aserGlyAla 120

seq_name: /SIDSI/gcgdata/geneseq/geneseqp-emb1/AA2000.DAT:AAV75271
seq_documentation_block:
ID AAV75271 standard; Protein; 119 AA.
XX
AC AAV75271;
XX
DT 21-MAR-2000 (first entry)
XX
DE Neisseria gonorrhoeae ORF 628 protein sequence SEQ ID NO:2016.
XX
KW Neisseria meningitidis; Neisseria gonorrhoeae; antigen; vaccine;
KW antigen; diagnosis; immunogenic; infection; meningitis; septicemia;
KW antibacterial; gene therapy.
XX
OS Neisseria gonorrhoeae.
XX
PN WO957280-A2.
XX
PD 11-NOV-1999.
XX
PF 30-APR-1999; 99WO-US09346.
XX
PR 01-MAY-1998; 98US-0083758.
PR 31-JUL-1998; 98US-0094869.
PR 02-SEP-1998; 98US-0098994.
PR 02-SEP-1998; 98US-0099062.
PR 09-OCT-1998; 98US-0103749.
PR 09-OCT-1998; 98US-0103794.
PR 09-OCT-1998; 98US-0103796.
PR 25-FEB-1999; 99US-0121528.
XX
PA (CHIR ) CHIRON CORP.
PA (GENO-) INST GENOMIC RES.
XX
PI Fraser C, Galeotti C, Grandi G, Hickey E, Maignani V, Mora M,
PI Petersen J, Piza M, Rappuoli R, Ratti G, Scalato E, Scarselli M,
PI Tettelin H, Venter JC;
XX
XX WPI: 2000-062150/05.
DR N-PSDB: AA254033.
XX
PT Novel Neisserial polypeptides predicted to be useful antigens for
PT vaccines and diagnostics
XX
PS Claim 2; Page 1003; 1453pp; English.
XX
XX AA253015 to AA254536, AA254577 to AA254615, and AAV74253 to AAV75941
XX represent novel Neisseria meningitidis and N. gonorrhoeae polynucleotides
XX and polypeptides. AA254537 to AA254576 and AA254616 to AA254473 represent
XX PCR primers used in the exemplification of the present invention. The
XX polypeptides, the polynucleotides, antibodies and compositions of
XX the invention can be used as vaccines, as diagnostic reagents, and as
XX immunogenic compositions. The polypeptides can be used in the
XX manufacture of medicaments for treating or preventing infection due to
XX Neisserial bacteria (e.g. meningitis and septicemia), to detect the
XX presence of Neisseria bacteria, or to raise antibodies. They may also
XX be used to screen for agonists or antagonists, which may themselves
XX have as antibacterial agents. The polynucleotides of the invention
XX may also be used in gene therapy protocols.
XX
SQ Sequence 119 AA;

```

```

alignment_scores:
  Quality: 487.50      Length: 119
  Ratio: 4.432
  Percent Similarity: 92.437      Percent Identity: 84.874

alignment_block:
US-09-303-518D-127/rev x AAV75271 ..

Align seg 1/1 to: AAV75271 from: 1 to: 119

674 ATGTGGCTGCACTCAACCGCGGATGCGGCGCGCGCAATTCATGTGT 625
|||||
1 MetCysValProLeuLysProAlaGlyCysGlyProProAsnSerCysVa 17
624 TTCGATGTGGCAGCATTTTCAGACGGCACGTCGTGGCGCCAGCTTCAC 575
|||||
17 IserIleLeuAlaAlaPheSerAspGlyThrSerAlaProAlaAlaLeuH 34
574 ACACATGGATTTTACGCTCGTCAACGGCTCAATACCAACANACNTCGT 525
|||||
34 IsthTrpIleLeuArgSerValArgArgLeuAsnThrAsnArgProArg 50
524 CTGAATCMTGNCGGCTCTTGTATCACACACAGGCTGCGCCNAG 475
|||||
51 LeuLysSerSerAlaAlaSerLeuMetThrValGlySerAlaAlaSe 67
474 CGGATTTGGTGTCCATCGCATTCGACGAAGATGGCAACGGCTGCGATCGA 425
|||||
67 rGlyLeuValSerIleAlaLeuThrLysMetAlaAsnGlySerAlaSerT 84
424 CGGACGAGATTTTCTGACGACGAGGNTACGACGCGAGTCACAAACCG 375
|||||
84 hrAlaGlyIleLeuLeuAsnGlyArgValArgSerAlaValHisLysPro 100
374 GATTGGATTCAGATTCGNNCANNANTTCNTGCCGCTTACGTTGGCAACGC 325
|||||
101 Asp...IleArgLeuArgArgThrPheSerLeuLeuAsnPhaAlaSerAl 116
324 TTCGGGGC 318
|||||
116 aserGly 118

seq_name: /SIDSI/gcgdata/geneseq/geneseqp-emb1/AA2000.DAT:AAV82082
seq_documentation_block:
ID AAV82082 standard; Protein; 467 AA.
XX
XX AAV82082;
XX
AC AAV82082;
XX
DT 01-JUN-2000 (first entry)
XX
DE Chlamydia pneumoniae antigen CPN100605 protein SEQ ID NO:2.
XX
XX Chlamydia pneumoniae; antigen; CPN100605 protein; immunisation;
XX vaccine; infection; antibacterial; antiinflammatory; bronchitis;
XX community acquired pneumonia; upper respiratory tract infection;
XX sinusitis.
XX
XX Chlamydia pneumoniae.
XX
XX WO200006742-A2.
XX
PD 10-FEB-2000.
XX
PF 27-JUL-1999; 99WO-IB01331.
XX
PF 27-JUL-1998; 98US-0094195.
XX
PR 26-JUL-1999; 99US-0361443.
XX
PA (CONN-) CONNUGHT LAB LTD.
XX

```

PI Murdin AD, Oomen RP;
 XX MPI; 2000-205466/18.
 DR N-PSDB; AA295378.
 XX

PT Chlamydia pneumoniae antigens used for immunization and protection
 against Chlamydia diseases -
 XX
 PS Claim 6; Fig 1; 48pp; English.

XX
 CC The present sequence represents the Chlamydia pneumoniae antigen
 CC CPN100605 protein. The CPN100605 protein has antibacterial and
 CC antiinflammatory activities. The Chlamydia pneumoniae CPN100605
 CC polynucleotide and protein can be used in vaccination methods for
 CC preventing and treating Chlamydia infection (e.g. infections caused by
 CC C. trachomatis, C. psittaci, C. pneumoniae or C. pecorum). The
 CC polynucleotide can be used to produce the protein recombinantly. In the
 CC construction of vaccine vectors, as a vaccine agent, and in the
 CC construction of an attenuated Chlamydia strain. The protein are also be
 CC useful as a vaccine agent, and for the preparation of medicaments for
 CC treating or preventing Chlamydia infection, e.g. community acquired
 CC pneumonia, and upper respiratory tract infections such as bronchitis and
 CC sinusitis.
 CC
 XX

SQ Sequence 467 AA:

alignment_scores:
 Quality: 447.50 Length: 468
 Ratio: 1.565 Gaps: 15
 Percent Similarity: 61.111 Percent Identity: 29.487

alignment_block:
 US-09-303-518d-127 x AAY82082 ..

Align seg 1/1 to: AAY82082 from: 1 to: 467

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4 ATTAATAATCAAAAAGGTCTAAACCTGCCCATCGCGGCGACACCGAGCA 53
||| : : : : : : : : : : : : : : : : : : : : : : : : : :
3 ILEthValAlsnArGILysLeuAspLeuSerLeuGlnGlySerProLysG1 19
: : : : : : : : : : : : : : : : : : : : : : : : : :
54 AGTCATTTATAGACGGCGCGCATTTACCGAAGTCGGCTTGCGGAGAG 103
: : : : : : : : : : : : : : : : : : : : : : : : : :
19 uSerGlyPheTyAlsnLysILeAspProLurPheValSerIle..... 33
104 AATATGCGGATATGCGCCCGC.....TNGATGAAGATC 135
||| : : : : : : : : : : : : : : : : : : : : : : : :
34 .....AspLeuArGProPheGlnProLeuSerLeuLysLeuVal 47
136 AAGGAAGCGGATCGCGTCAAAAAGGCAAGTGTGTTGAAGACAAAAA 185
: : : : : : : : : : : : : : : : : : : : : : : : : :
48 GILuGlnGlyAspAlaValLysSerGlyAlaProIleAlaGluTyrLysH1 64
186 GNAATCGGCGGTGGTGTATACCGCGCCGCTTCAAGCAAAATCGCGGCA 235
||| : : : : : : : : : : : : : : : : : : : : : : : :
64 sPheProAsnIleTyrIleThrSerHisValSerGlyValValThrAlaI 81
236 TCCATCGCGCGGAGAAAAGCGCTACTTCAAGTCGGTGTGATTT...GCCGTT 282
||| : : : : : : : : : : : : : : : : : : : : : : : :
81 IeArGArGILysILsnLysArGSerLeuLeuAspValIleIleLysLysThr 97
283 GAAGGCAACGACGAATGAGTTCGACGCTACGCGCCGCAAGCGTGTGC 332
||| : : : : : : : : : : : : : : : : : : : : : : : :
98 ProGILyProThrSerThrGluTyr.....ThrTyrAspLeuG1 110
333 AACTTAAGCGGCGGANGAANTNNGCAATCTGATCCAAATCGGTTGT 382
: : : : : : : : : : : : : : : : : : : : : : : : : :
110 nThrLeuSerArGSerArPheSerGILuIlePheLysGILuSnGlyLeuP 127
383 GGACGCGCGTGGTANCGTCCGTTCCACAAAATCCCTGCGCGTGATGCC 432
||| : : : : : : : : : : : : : : : : : : : : : : : :
127 heAlaLeuIleLysGILuArGProPheAsp...IleProAlaIleProThr 142

```

```

433 GAG...CGGTGGCATCTTTCGTAATGCGATGACACCAATCCGCTGNC 479
: : : : : : : : : : : : : : : : : : : : : : : : : :
143 GlnThrProArGAspValPheILeAsnLeuAlaAspAsnArGProPheTh 159
480 GGCAGACCGCTGTGGTTGTGATC.....AAAGAAGCCG 511
: : : : : : : : : : : : : : : : : : : : : : : : : :
159 rProSerProGILuLysILsnLeuAlaLeuPheSerArGILuGluGlyP 176
512 NCGANGATTTTCAGACGANGTGTGTATGAGCGGCTTGACGAGCGT 561
||| : : : : : : : : : : : : : : : : : : : : : : : :
176 heTyValPheValValGlyValArGAlaIleAlaLysLeuPheLysLeu 192
562 AAAATCCAGTGTGTGAAGGCACTGGCCGACACGTCGCGTGAATGCG 611
: : : : : : : : : : : : : : : : : : : : : : : : : :
193 ArGProILsnIleValPheArGAspArGLeuThrLeuProThrGlnGlu 209
612 TCCCAACATC...GAACACATGAATTTGGCGCGCGCATCGCGCGGT 658
||| : : : : : : : : : : : : : : : : : : : : : : : :
209 uLysThrIleAlaILsnILsnILsnILsnILsnILsnILsnILsnILsn 226
659 TGAAGGCGACGACATTTGATTTGATTCATGACCGCGTGTGCAACAA... 705
: : : : : : : : : : : : : : : : : : : : : : : : : :
226 ePProSerILsnILsnILsnILsnILsnILsnILsnILsnILsnILsn 242
706 ACCGTTTGACACATCATATATATCAAGATGATATGCAATCGGACGCT 755
: : : : : : : : : : : : : : : : : : : : : : : : : :
243 ValValPheThrLeuSerPheGlnAspValLeuThrILeGILuILsnLeuP 259
756 TGCACACGCGCGTGTGAACACGACGCGCGATGCTTGGGTGGTTCG 805
||| : : : : : : : : : : : : : : : : : : : : : : : :
259 eLeuLysGILuArGILeILsnILsnILsnILsnILsnILsnILsnILsn 276
806 AAGTCACAAACCA...CGCCTTTGGCTACCGTTTGGTGGCGA 849
: : : : : : : : : : : : : : : : : : : : : : : : : :
276 IeLeuLysSerSerLeuArGArGILuValIleThrILsnLysGILuLaser 292
850 GTATCGCAA...ATTACTGCGCGGCAATGGTGTGATCCGACGACCGCGT 896
||| : : : : : : : : : : : : : : : : : : : : : : : :
293 PheSerSerLeuILeAsnILeAsnILsnArPheILserILsnILsnILsn 308
897 GATTTTCGGTTCGATTTGAACGCGCGGATTTACACAGCGCGCACGAT 945
: : : : : : : : : : : : : : : : : : : : : : : : : :
308 uILeSerGILuAspProILsnILsnILsnILsnILsnILsnILsnILsn 325
946 ..TATTTGGACGCTACACATGATTCGATTCGATTCGAAAGCGCGC 993
||| : : : : : : : : : : : : : : : : : : : : : : : :
325 rOrPheLeuGILuPheArGAspILsnILsnILsnILsnILsnILsnILsn 341
994 AGCAAAAGCTGTTCGCTGAGTTCGCGCGACGCGGACAAATATCCAT 1043
: : : : : : : : : : : : : : : : : : : : : : : : : :
342 LysArGILuLeuPheSerPheLeuArGILeGILuPheAsnLysProThrP 358
1044 CAGCGGTACGACGCGCTGCGCATTTCTGAAAACAAATCTTCAAGITCA 1093
||| : : : : : : : : : : : : : : : : : : : : : : : :
358 eThrLysThrTyrLeuSerGILuPhePheLysLysLysArG...ThrTyr 374
1094 CG.....ACAGCGCTACGCGGTGCGCGACGCGGCAAGATGCGCGAT 1134
||| : : : : : : : : : : : : : : : : : : : : : : : :
374 hAsnProArGILuPheILsnILsnILsnILsnILsnILsnILsnILsn 390
1135 GGTACTTACAGCGCGTAAATGCGCTGACATCTTCCTACGCTGCTTT 1184
||| : : : : : : : : : : : : : : : : : : : : : : : :
391 AspILeTyrAspLysValMetProMetArGILeProValILsnProLeuI 407
1185 GCGCGATTTATGCTGCGGATACGACGACGCGGCAAGATGCGGTGCT 1234
: : : : : : : : : : : : : : : : : : : : : : : : : :
407 eLysAlaValILeThrLysILsnPheAspLeuILsnGILuLeuGILuPheL 424
1235 TGGAAATTTGACGAAGACATCGCTTTTGACAGCTTCGCTGCGCGGCG 1284
||| : : : : : : : : : : : : : : : : : : : : : : : :
424 euILuValCysGILuAspPheILsnILsnILsnILsnILsnILsnILsn 440
1285 AATATCAATATANGCGCGCTGTTCGTAAGTGTGAAACCGTTGAGAA 1334

```

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||| |||
441 LysThrGluMetLeuThrIleValIleGluSerLeuIleGluTyrAlaIy 457
1335 GGA 1338
|||
457 GGU 458

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seq_name: /SIDSI/gcgdata/geneseq/geneseq-emb1/AA1999.DAT:AAV35375

seq_documentation_block:

ID AAV35375 standard; Protein; 469 AA.

AAV35375;

13-SEP-1999 (first entry)

Chlamydia pneumoniae transmembrane protein sequence.

Respiratory disease; pneumonia; bronchitis; heart disease; sarcoidosis;

sinusitis; purulent otitis media; erythema nodosum; pharyngitis;

vacine; neutralising epitope.

Chlamydia pneumoniae.

W09927105-A2.

03-JUN-1999.

20-NOV-1998; 98WO-1B01890.

04-NOV-1998; 98US-0107078.

21-NOV-1997; 97FR-0014673.

(GEST) GENSET.

Griffais R;

WPI: 1999-357842/30.

Genome sequence of Chlamydia pneumoniae

Page 1170-1171; Disclosure: 1912pp; English.

AAV34584-Y35879 represent the proteins encoded by all the open reading frames in the complete genome (see AAX91990) of Chlamydia pneumoniae. C. pneumoniae causes respiratory disease such as pneumonia and bronchitis and is thought to be a contributing factor in heart disease, sarcoidosis, sinusitis, purulent otitis media, erythema nodosum or pharyngitis. The polypeptides encoded by the open reading frames of the C. pneumoniae genome (see AAV34584-Y35879) can be used in immunogenic compositions as vaccines. Vectors containing C. pneumoniae nucleotide sequences can also be used as immunogenic compositions, especially where the vector directs the expression of a neutralising epitope of C. pneumoniae.

Sequence 469 AA:

alignment_scores:
 quality: 447.50 Length: 468
 Ratio: 1.565 Gaps: 15
 Percent Similarity: 61.111 Percent Identity: 29.487

alignment_block:
 US-09-303-518D-127 x AAV35375 ..

Align seg 1/1 to: AAV35375 from: 1 to: 469

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4 ATTAATAATCAAAAAGGTCTTAACCTGCCATCGGGGAGACCGGAGCA 53
||| :|||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
5 IleThrValAsnArgGlyLeuAspLeuSerLeuIleGlnGlySerProIysG 21
54 AGTCATTATGACGGCCCGCTCATTCACGAAGTCGCGTTGCTGGCAG 103

```

```

21 userGlyPheTyrAsnIleValIleAspProGluPheValSerIle..... 35
104 AATATGCCGGTATGCCCC.....TNGATGAAGTC 135
36 .....AspLeuArgProPheGlnProLeuSerLeuIleValIle 49
136 AAGGAAGCGCATGCGTCAAAAAGGCCAAGTCTGTTTGAAGACAAA 185
50 GlnGlnGlyAspAlaValIleCysSerGlyAlaProIleAlaGlnGlySer 66
186 GNAATCCGGGGCGGTGTTTACCGCGCCGCTTTCAGCAAAATCGCCGCCA 235
66 sPheProAsnThrTyrIleThrSerHisValSerGlyValIleThrAla 83
236 TCCATCGCGCGCAAAAGCGCTACTTCAGCGGTGCGTGAAT...GCCGT 282
83 leArgArgGlyAsnIleValIleValIleValIleValIleValIleVal 99
283 GAAGCAGCAGCAAAATCGATTCGAACGCTACGCCCGCCGAGCGTTGGC 332
100 ProGlyProThrSerThrGluTyr.....ThrTyrAspLeuGln 112
333 AAACCTTAAGCGCGGANGANTNNGNCAATTCGATCCATCCGTTGT 382
112 nThrLeuSerArgSerAspLeuSerGluIlePheIleValIleValIleVal 129
383 GGAATCGCGCGGTTCGTCGTCGTCGTCGTCGTCGTCGTCGTCGTCGTCG 432
129 heAlaLeuIleValIleValIleValIleValIleValIleValIleVal 144
433 GAG...CCGTTGCCATCTTCGTCGTCGTCGTCGTCGTCGTCGTCGTCG 479
145 GlnThrProArgAspValIlePheIleAsnLeuAlaAspAsnArgProPhe 161
480 GCGAGACCCCTGTTGATC.....AAGAGCCG 511
161 rProSerProGluIleValIleValIleValIleValIleValIleValIle 178
512 NCGANGATTTCAGACGANGTGTGATTCGATTCGATTCGATTCGATTCGAT 561
178 heTyrValPheValIleValIleValIleValIleValIleValIleVal 194
562 AAATCCATGTGTGTAAGCAGTGGCGCAGACGTCGCGTCAAAATGC 611
195 ArgProHisIleValIlePheArgAspArgLeuThrIleThrIleGlnGlu 211
612 TGCCACATC...GAACACATGAATTCGCGCGCGCGCGCGCGCGGT 658
211 ulsThrIleAlaHisIleHisIleHisIleHisIleHisIleHisIleHis 228
659 TGAGTGCAGCGACATTCATTCATTCATTCATTCATTCATTCATTCATTC 705
228 erProSerIleHisIleHisIleHisIleHisIleHisIleHisIleHis 244
706 ACCGTTTGACCATCATTCATTCATTCATTCATTCATTCATTCATTCATTC 755
245 ValIleValIleThrLeuSerPheGlnAspValIleThrIleHisIleHis 261
756 TGCAACAGCGCGCTGTAACACCGAGCGGTGATTCGTTGGGTGCTC 805
261 eleuIleGlyArgIleIleHisIleGlnValIleThrIleAlaIleGlyThr 278
806 AAGTCACAAACCA.....GCGCTCTGCGTACCGTTTGGTGGCGAAA 849
278 leuIleSerSerLeuArgArgTyrValIleThrIleHisIleHisIleHis 294
850 GATATGCA...ATTACGCGCGCGATGATGTTGTTGTTGTTGTTGTTGTT 896
295 PheSerSerLeuIleHisIleHisIleHisIleHisIleHisIleHisIle 310
897 GATTTCGCGTTCGATTCGATTCGATTCGATTCGATTCGATTCGATTCGAT 945

```



```

181 aLgIyAlaGlAlaIleAlaLysLeuPheGlyLeuLysProHisIleIle 197
577 AAGGCGAGTGGGCGAGACGTCGCTGTAATGCTGCACATC...GA 623
198 SerThrAspArgLeuThrLeuProThrGlnAspLeuValSerIleAlaHi 214
624 AACACATGAATTCGGGCGCCGATCCGGCGGCTTGAATGACGACGACA 673
214 sleuHisThrIleAspGlyProPheProSerGlySerProSerThrHisI 231
674 TTGATTTCAATGAGCGGTC...GGTCAACAAACAAACGTTGGACCATC 720
231 IeHisHisIleAlaArgIleArgAsnGluArgAspAlaIlePheThrIle 247
721 AATTATCAAGATGTAATGATGCATGAGCGCTTTGTAACAGCGCCCTCT 770
248 SerPheGlnGluValLeuSerIleGlyHisLeuPheLeuGlyPheVa 264
771 GAACACCGAGCGCGTATGCTTGGGTGCTCAAGTC.....ACA 814
264 IleuGlyGlnGlnIleValAlaLeuAlaGlySerAlaLeuProProSerG 281
815 AACCGCGCTTCGCTACCGCTTGGGCGAAGATGATGCAATTAATCT 864
281 IuArgIlystyLeuIleThrAlaLysGlyAlaSerPheSerAspLeu 297
865 GCGGCGCAATTGGTGTGACGAGCAGAC...CGCGTATTTCCGTTCCGT 911
298 ProLysAspIlePheSerSerAspGluIleThrLeuIleSerGlySerP 912
912 ATTGACGCGCGCATTCACACAGCGCGCGCAT...TATTGGAGCGCT 958
314 oLeuThrGlyArgLeuLysIlyGluGlnAsnProCysLeuGlyMeta 331
959 ACCACATCAAGATTTCCGTTATGCAAGAGCGCGCAGACAAAGCTGTT 1008
331 rGaAspHisThrIleThrLeuLeuProAsnProLysThrArgGluSerP 347
1009 GCGTGGGTGGCGCGCGACGACAAATACTCCATCAGCGGTACGACCT 1058
348 SerPheLeuArgLeuGlyTyrPasnIlyLeuThrValThrArgThrIly 364
1059 CGGCGATTTCTGAAAAACAA.....CTCTTCAAGTTACGACGACCG 1102
364 uSerGlyPhePheLysArgIlyArgValPheMetAspMetAspThrsm 381
1103 TCAACGCGTGGCGCGCGCATGCTGCGCATGTGCTACTTACGAGCGCT 1152
381 eThIsGlyGluLysArgProIleIleAspAlaGluIleTyrGluArgVal 397
1153 ATGCCGCTAGACATCCCTGCTACCCCTGTTGGCGCATTTAATCGTCG 1202
398 SerAlaIleProValProValAlaLeuIleIleLysAlaLeuGluThrG 414
1203 CGATACCGACAGCGCGACATTTGGTGTGTAATGGACGACAGAG 1252
414 nasnheGluGluAlaLysArgLeuGlyLeuLeuGluValAlaProGlu 431
1253 ACCCTGCTTTGACGCTTCTGCTGCGCGCGCAATACGATTAAGGCCG 1302
431 spPheAlaLeuProThrPheIleAspProSerLysThrGluMetPheSer 447
1303 CTGTTCGTAAGGTGCTGTAACCTTGAAGA 1335
448 IleValIlyGlnSerLeuLeuArgThrGlnLys 458
seq_name: /SIDS1/gcgdata/geneseq/geneseq-emb1/AA1999.DAT:AAV34467
seq_documentation_block:
ID AAV34467 standard; Protein; 443 AA.
XX
AC AAV34467;

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XX
DT 25-AUG-1999 (first entry)
XX
DE Porphyromonas gingivalis protein PG122.
XX
KW Porphyromonas gingivalis; PG; periodontal disease; gingivitis;
XX vaccine; antigenic.
XX
OS Porphyromonas gingivalis.
XX
PN WO929870-A1.
XX
PD 17-JUN-1999.
XX
PF 10-DEC-1998; 98WC-AU01023.
XX
PR 04-AUG-1998; 98AU-0005028.
PR 10-DEC-1997; 97AU-0000839.
PR 31-DEC-1997; 97AU-0001182.
PR 30-JAN-1998; 98AU-0001546.
PR 10-MAR-1998; 98AU-0002264.
PR 09-APR-1998; 98AU-0002911.
PR 23-APR-1998; 98AU-0003128.
PR 05-MAY-1998; 98AU-0003338.
PR 22-MAY-1998; 98AU-0003654.
PR 29-JUL-1998; 98AU-0004917.
XX
PA (CSLC-) CSL LTD.
XX
PI Agius CT, Barr IG, Hocking DM, Margetts MB, Patterson MA;
PI Ross BC, Roedel LJ, Webb EA;
XX
DR WPI; 1999-385613/32.
XX
PT N-PSDB; AAX91685.
XX
PF Antigenic Porphyromonas gingivalis peptides for preventing
PF gingivitis
XX
PS Claim 1; Page 445-446; 588bp; English.
XX
CC AAX91536 to AAX91801 encode two hundred and sixty six antigenic
CC Porphyromonas gingivalis (PG) polypeptide sequences given in AAY34318 to
CC AAY34363. AAX91802 to AAX91989 represent PCR primers used in the
CC isolation of the PG polypeptides. The PG polypeptides have antibacterial
CC activity with a vaccine mechanism of action. The PG polypeptides can be
CC used as vaccines especially against Porphyromonas gingivalis. Probes can
CC be used to detect Porphyromonas gingivalis in standard hybridisation
CC assays. Porphyromonas gingivalis is involved in periodontal disease
CC especially gingivitis.
XX
SQ Sequence 443 AA;

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alignment_scores:

Quality:	159.50	Length:	469
Ratio:	0.697	Gaps:	17
Percent Similarity:	48.827	Percent Identity:	19.616

alignment_block:

US-09-303-518D-127 x AAV34467 ..

Align seg 1/1 to: AAV34467 from: 1 to: 443

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37 GCGGCGAGACCGAGACGATTCATTTGACGGCGCCGCTATTCGCAAGT 86
|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
19 AlaglyysProValGluValleu.....ProIleProSerGlnVa 32
87 CGCGTCTCTGGCGAGATATGCGCGTATGCGCCCTTGATGTAAGTCA 136
|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||
32 IValIleProLeuGlyGlnHisIleGlyAlaProAlaThrAlaThrVal 49
137 AGNAGCGATGCCGTCACAAAAGGCCAGCTGCTGTTGAAGACAAAG 186
|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||

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49 ystyscltyspgluvallysvalglythrlellelaclnlaaglygly 65
187 NATCGGCGGTGGTGTACCGCGCGGTTTACGCAAAATCGCGCCAT 236
66 PhevalSerAlaasnIleHisSerValSerGlyLysValLeuLysI 82
237 C.....CATCGCGCGCAAAAGCGCTACTTCACTGGTGGTGGAT 277
82 eapaaenValTyrAspSerSerGlyTyrProLysProAlaValPheI 99
278 CGGTGAAGCAAGCAAGCAATCGAGTTCGAACGCTACCGCCCAAGCG 327
99 erValGlucltyspgluutpugluGlyIleAspArgSerProAlaI 115
328 TTG.....GCAAACTTAAGCGCGCGANGAANTNGNNGCAATCTGAT 371
116 ValLysGluCysasnLeuaspAlaLysGluIleValAlaLysIle 132
372 ATCCGGTTTGTGACTGCGCTCGGTANCCGCTTC..... 408
132 aaIaGlyIle..ValGlyLeuGlyGlyAlaThrPheProThrHisVal 148
409 .....AGCAAAATCCCTGCGCTGATGCGCGAGCGGTGCAATCTGCT 453
148 yLeuSerProProProGlyasnLysAlaGlu.....IleLeuIle 162
454 AATGGATGGACCAATCCG...CTNGCGCGAGACCTGTGTGTAT 500
163 AsnAlaValGluCysGluProTyrLeuThrSerAspHisValLeuMe 179
501 CAAGACCGCGCGANGATTCACAGCANGTNGCTGTATGAGCGGT 550
179 uGluHisGlyGluGluIleMetIleGlyValSerIleLeuMetLysAl 196
551 TGACCGACCGTAA..... 564
196 leGluValasnLysAlaValIleGlyValGluasnAsnLysLysAsp 212
565 ...ATCCATGTGTGAAGGACGCTGGCGCGAGCGTGCCTGTGAAATGC 611
213 IleAlaHisLeuThrLysLeuAlaThrAla.....Ty 223
612 TGCCACATCGAACAACATGATTCGGCGCGCGCATCCGCGGTGA 661
223 rProGlyIleGluValMetProLeuLysValGluTyrProGluGly 240
662 GTGCACGCGACATTCATTTCATT..... 684
240 IuLysGluLeuIleaspAlaValIleArgLysGluValLysSerGly 256
685 GAGCGCGTGGTGAACAAACACCGTTGACCATCAATTATCAAGTGT 734
257 LeuProIleSerThrGlyAlaValValGln.....AsnValGlyThr 271
735 AATGCGATCGAAGCTTTGTTGCAACAGCGCTGACACCGGCGCG 784
271 lPheAlaValTyrGluValValGlnLysasnLysProLeuValGlu 288
785 TGATGTGGTGGTGGTTCACAGTCAACAACACCGCTTTCGCTAC 834
288 leValThrValThrGlyLysLysLeuSerArgProSerAsnLeuLeu 304
835 GTTTGGTGGCGAAGATATGCAAAAT.....ACTGCGCGCAATTTGT 878
305 ArgIleGlyThrProIleAlaAlaLeuIleGluAlaGlyLeuLeu 321
321 ogIuasnThrGlyLysIleIleGlyGly..... 331
929 CACAAGCGCGCGACGATTAATTGGGCGCTACACCAATAGATTCCGT 978
331 ..... 331

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979 ATCGAAGAGCGCCAGCAAGACCTGTGCGTGGGTGGCGCGCACCC 1028
332 .....PrometMetGlyArgAlaLeuLeuSer 341
1029 GGACAAATACATCCATCAGCGGTACGACCTTCGGCATTTCTGAAA 1078
341 oASP...ValProValThrLysGlySerSerGlyValLeuIleAsp 357
1079 AACTCTTCAAGTTCCAGACAGCCGTCACGCGTGCAGCGCGCATG 1128
357 rg.....GluGluAlaValAlaArgLysPrometArgAspCysIle 369
1129 CCGATTGTGATACGAGCGCGTAATGCGCTGACATCTGCTACCT 1178
370 ArgCysAlaLysCysValGlyValLysPrometGlyLeuasnPro 386
1179 GCTTTGCGCGATTTATGCTGCGCGATACCGAGCGCGCA..... 1221
386 eleuMetArgAspThrLeuTyrLysSerThrPgluThrAlaGluLys 403
1222 ..GCATTGGGTTCTTGAATTGGACAGAGACCTCGCTTGTGAC 1269
403 snValAlaAspCysIleGlyCysGlySer.....CysSer 414
1270 TTGCTGTGCGCGCGCAATACGATANGCGCGCTTGCATGAGTGT 1319
415 PheThrCysProAlaasnArgProLeuLeuAspTyrIleArgGlnAl 431
1320 GGAACCC 1326
431 sLysThr 433

seq_name: /SIDSL/gcgdata/geneseq/geneseq-emb1/AA1999.DAT:AAV34343
seq_documentation_block:
ID AAV34343 standard; Protein: 451 AA.
AC AAV34343;
DE 25-AUG-1999 (first entry)
XX Porphyromonas gingivalis protein PG122.
XX DE Porphyromonas gingivalis; PG, periodontal disease; gingivitis;
XX KW Porphyromonas gingivalis;
XX KW vaccine; antigenic.
XX OS Porphyromonas gingivalis.
XX PN WO9929870-A1.
XX PD 17-JUN-1999.
XX PE 10-DEC-1998; 98WO-AU01023.
XX PR 04-AUG-1998; 98AU-0005028.
XX PR 10-DEC-1997; 97AU-0000839.
XX PR 31-DEC-1997; 97AU-0001182.
XX PR 30-JAN-1998; 98AU-0001546.
XX PR 10-MAR-1998; 98AU-0002264.
XX PR 09-APR-1998; 98AU-0002911.
XX PR 23-APR-1998; 98AU-0003128.
XX PR 05-MAY-1998; 98AU-000338.
XX PR 22-MAY-1998; 98AU-0003654.
XX PR 29-JUL-1998; 98AU-0004917.
XX PA (CSLC-) CSL LTD.
XX PI Agius CT, Barr IG, Hocking DM, Margetts MB, Patterson MA;
XX PI Ross BC, Rothel LJ, Webb EA;
XX WPI; 1999-385613/32.
XX DR N-PSDB; AAX91561.

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541 rSerGlyArgArgSerGlyAlaThrThrMetIleThrProThrValSerS 558
1053 GACCTGCGGCGCTTCTGAAAACAACTCTCAAGTTCAGACGACCG 1102
558 erProAlaSerThrArgLysSerSerAlaLysCysAlaArgSerPro 574
1103 TCACGCGTGGCGACCGCGCATGTGCGCATGTGCTACTTACGACGCGGTA 1152
575 ThrThrIleValIValArgSer...CysArgIleValIArgIleSerAla.. 589
1153 ATGCGCGTAGACATCGCTACCTGCTGCTTGGCGCATTTAATCGTCGG 1202
590 .....CysThrTrpLysSerValA 596
1203 CGATACCGACGCGCGCAAGCATTTGGTCTGCTGGAATGGACGACAGAG 1252
596 rGAlaArgAlaProArgArg..... 602
1253 ACCTGCGCTTGGCGACGCTTGGTGGCGGCAATACGATANGGCC.. 1300
603 .....ThrGlyAlaSerGlyValLysArgIleThrAlaAlaPh 616
1301 .....CGCTGTGCGGTAA 1313
616 eleuProThrTrpGluProThrArgArgGlyArgArgArgCysCysAlaA 633
1314 GGTGC 1318
633 rGcys 634

seq_name: /SISL/gcgdata/geneseq/geneseq-emb1/AA2001.DAT:AA59826
seq_documentation_block:
ID AA59826 standard; Protein: 1615 AA.
AC AA59826;
DT 04-APR-2001 (first entry)
DE Protein #3 encoded by Tufd/E gene.
DE
DE Toluene degradation; enzyme; waste degradation; Tufd; Tufd.
KM
OS Thauera aromatica.
OS Xanthomonas maltophilia.
OS Geobacter metallireducens.
OS Azorhizobium toluyticus.
XX
XX WO200072650-A2.
XX
XX 07-DEC-2000.
XX
XX 24-MAY-2000; 2000WO-US14298.
XX
XX 01-JUN-1999; 99US-0323872.
XX
XX (UHOH-) UNITV OHIO.
XX
XX Coschigano PW;
XX
XX MPI: 2001-041080/05.
XX
XX N-PSDB: AAF23627.
XX
XX Composition comprising toluene degrading enzyme useful for biological
XX treatment of organic compounds, especially for degrading toluene or its
XX analogs
XX
XX Disclosure; Fig 12; 12pp; English.
XX
XX The present invention relates to toluene degrading enzyme genes and
XX proteins tufd (see AAF23629 and AAB59831), tufE (AAF23630 and AAB59832),
XX tufF (AAF23631 and AAB59833) and tufG (AAF23632 and AAB59834). The
XX toluene degrading enzymes are homologues of pyruvate formate lyase. The

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CC toluene degrading enzymes are useful for biological treatment of organic
CC compounds and in particular for the degradation of toluene and its
CC analogs contained in liquid or solid waste source. The present sequence
CC is a protein sequence encoded by toluene degrading enzyme gene, Tufd/E.
XX
XX Sequence 1615 AA;

alignment_scores:
Quality: 134.50 Length: 469
Ratio: 0.659 Gaps: 22
Percent Similarity: 43.497 Percent Identity: 22.175

alignment_block:
US-09-303-518D-127 x AA59826 ..
Align seg 1/1 to: AA59826 from: 1 to: 1615

35 TCAGCGGCGACGCGGACGAGCATGATTTATGACGGCGCGCTATTACCGAA 84
858 SerArgAlaProThrAlaLysSer.....SerAr 867
85 GTGCGCTTGGTGGCGGAGAAATATGCGGTATGCGCGCGCTATGAAAGT 134
867 gGlyArgThrIleCysSerSerSerProSerAlaAlaProThrProArgA 884
135 CAAGG..... 139
884 laArgThrProAlaThrThrProThrProSerSerArgIleProSerGly 900
140 AAGCGCATGCGCTGCAAAAAGCCAGTGTGTTGAGCAAAAAGNAT 189
901 SerAlaArgProSerProProSerSerSerAlaIleProArgArgThrAl 917
190 CCGGCGGTGTGTTTACCGCGCGCGGTTCAGGCAAAATGCGCGCATCCA 239
917 aArgArgArgCys.....AlaGlyPheSerSerAlaSerAlaThrAspS 932
240 TCAGCGGCAAAAGCGCGTACTTACGTGCGTGTGATTCGCGTGAAGCA 289
932 eAlaIleArgArgSerSerThrThrArgSerAlaArgSerArgAsn 948
290 ACGAGAAATGAGTTCGACGCTACGCGCGCGGAGCGTTGGCAACTTA 339
949 ThrProSerSerAlaSerThrAlaThrAlaProProThr..... 961
340 AGCGGCGANGAANNNGNNGCAATCGATTCGATCCGTTGTGTGACTGC 389
962 .....ArgLysProThrThrLysSerT 969
390 GCTGCGTANCCGCTCGTTCAGCAAAATCCGCGCGTGAATGCCAGCGGT 439
969 hrCysCysAlaCysArgProAlaSerThrVal..... 979
440 TCGCATCTTCGTCAATCGCATGACACCAATCCGTTGCGGCGAGACCT 489
980 .....AlaAlaArgArgLysLysProValArg..... 988
490 GTGTTTGATCAAGAGCGCGANGATTTGACAGCANGTNGTGGT 539
989 .....LysValAlaAlaGlnSerSerArgProSerGlyStrp. 1000
540 ATTGAGCGTTTGACCGAGCGTAATCATGTGTGAAGCGAGCTGCGG 589
1001 .....LysSerArgSer 1004
590 CAGACGTGCGCTGTAATAATGCTGCCACATCGAACAACATGATGCGC 639
1005 MetThrAlaThrThrGlyArgThrProThr.....CysAsnSerAl 1018
640 GCGCGCATGCGCGCGGTTGAGTGCGACGACGATTCATTCATTCAGCC 689
1018 aArg.....ArgProValIleSerThrArgArgSerProSerArgMet.Phe 1032

```

```

690 GGTGGTGCACAAACCGTTGGACATCATTAATGATGATG 739
1033 GlyArgLeuSerAlaSerSerIleAsnMetArgSerThrValSerAl 1049
740 CCATGGAGCGTTGTTGCACAGCGCGTGCACACCGCGCGTGAAT 789
1049 aProkArgThr.....CysArgAlaThrSerSerAlaSerCysArgC 1064
790 GCTTGGGTGGTTCTCAAGTCAACAAACCGCTTGGCTACCGTTT 839
1064 ys.....LeuSerCysProGlnSerThrThrAlaAlaTrpAsnSer 1077
840 GGG.....TGCAGAAAGTATGCGAAATTACTGCGGCGAAT 874
1078 GlyTrpThrProAlaProCysProSer...SerProMetAlaGlyThrThr 1093
875 TGGTTGACGACAGACACCGCGTGTTCGGTTCGGTATTGACGCGCG 924
1093 rArgSerArgArgSerSerArgArgThrProSerTrpProSerArgAsnT 1110
925 ATTACACAGGCGCGCACGATTTATTGGACGCTACCAATCAGATTTC 974
1110 rPyrSerArgArgArg.....AsnThrProSerSerAsnSer 1122
975 CGTTATCGAAGAGAGCC.....GCAGCAAGAGCTGT.....TC 1008
1123 AlAlaSerArgArgArgThrGlyLysValSerArgLysCysAlaSerThrSe 1139
1009 GGCTGGGTTCGCGCGACCGCGGACAAAT.....ACTCATTCACGGGTAC 1052
1139 rSerGlyArgArgSerGlyAlaThrThrMetIleThrProThrValSerS 1156
1053 GACCCGTGGCCATTCTCTGAAAACAACTCTCAAGTTCACGACAGCGC 1102
1156 erProAlaSerThrArgLysSerSerAlaAlaLysCysAlaArgSerPro 1172
1103 TCAACGGTGGCGACCGCGCATGTGTGTACTTACGACGCGCGTA 1152
1173 ThrThrLeuValValAlaArgSer...CysArgLeuValAlaArgLeuSerAla.. 1187
1153 ATGCCCGTACACATCTGCTACCTGCTTGGCGGATTTAATCGTCGG 1202
1188 .....CysThrTrpLysSerValAla 1194
1203 CGATACCGACAGCGCGCAAGCATGTGGTGTGATGATGACAGGAAG 1252
1194 rGAlaArgAlaProArgArg..... 1200
1253 ACCTCGCTTGTGCACCTTCGTCTGCGCGCAATATGCAATANGGCC.. 1300
1201 .....ThrGlyAlaSerGlyValLysArgGlnThrThrAlaAlaPh 1214
1301 .....CGCTGTTCGCTAA 1313
1214 eleuProThrTrpGluProThrArgArgGlyArgArgArgCysAlaAla 1231
1314 GGTGC 1318
1231 rGCys 1232
seq_name: /SIDS1/gcdata/geneseq/geneseq-emb1/AA2001.DAT: AAB59817
seq_documentation_block:
ID AAB59817 standard; Protein; 999 AA.
XX
AC AAB59817;
XX
DT 04-APR-2001 (first entry)
XX
DE Tuta protein #8.
XX
KW Toluene degradation; enzyme; waste degradation; Tuta.

```

```

XX Thauera aromatica.
OS Xanthomonas maltophilia.
OS Geobacter metallireducens.
OS Azarcus toluyticus.
PN MO200072650-A2.
PD 07-DEC-2000.
PF 24-MAY-2000; 2000WO-US14298.
PR 01-JUN-1999; 99US-0323872.
PA (UTOH-) UNIV OHIO.
PI Coschignano PW.
DR WPI: 2001-041080/05.
DR N-PSDB: AAF23625, AAF23627.
XX
XX Composition comprising toluene degrading enzyme useful for biological
XX treatment of organic compounds, especially for degrading toluene or its
XX analogs
XX
XX Disclosure; Fig 5; 122pp; English.
XX
XX The present invention relates to toluene degrading enzyme genes and
XX proteins tuta (see AAF23629 and AAB59831), tuta (AAF23630 and AAB59832),
XX tuta (AAF23631 and AAB59833) and tuta (AAF23632 and AAB59834). The
XX toluene degrading enzymes are homologues of pyruvate formate lyase. The
XX toluene degrading enzymes are useful for biological treatment of organic
XX compounds and in particular for the degradation of toluene and its
XX analogs contained in liquid or solid waste source. The present sequence
XX is a protein sequence for toluene degrading enzyme, Tuta.
XX
XX Sequence 999 AA:
XX
XX alignment_scores:
XX Quality: 132.00 Length: 504
XX Ratio: 0.706 Gaps: 25
XX Percent Similarity: 37.103 Percent Identity: 22.619
XX
XX alignment_block:
XX US-09-303-518D-127 x AAB59817 ..
XX
XX Align seg 1/1 to: AAB59817 from: 1 to: 999
XX
XX 81 CGAAGTGGCTTGTGTCGCAAGATATGCGGATGCGCCCTNGATGA 130
XX ||||| ||| ||| ||||| ||||| ||||| ||||| ||||| |||||
XX 117 ArgSerArgCysSerProAspArgCysArgTrpSerArgCysSerS 133
XX 131 AAGTCAAGAGAGCGATGCGCTCAAAAAGCCCAAGTGTGTTGAAGC 180
XX : : : : : ||||| ||||| ||||| ||||| ||||| |||||
XX 133 rProSerProArgArgCysProProSerSerProAlaGly..... 146
XX 181 AAAAAGNATCCGGGCGTGTGTTACCGCGCGCGTTCAGCAAAATGCC 230
XX : : : : : ||| ||| ||| ||||| ||||| ||||| |||||
XX 147 .....AlaProGlyAlaThrCysSerArgArgProPheserAlaSerArg 161
XX 231 CGCCATTCATCGCGCGAAGAGCGCTACTTCAAGTGTGCTGAT...TG 277
XX : : : : : ||| ||||| ||||| ||||| ||||| |||||
XX 162 AspSerAlaGlyProArgAlaAlaArgPheArgArgCysArgAspAla 178
XX 278 CGCTGAAGGCAACGA..... 293
XX 178 sGluArgArgAlaArgCysProGlyProArgSerAlaProSerIleArg 195
XX 294 .....CGAATCGAGTTGACAGCTACGCGCGCGCGAAGCGTTGGCAA 335
XX ||||| ||| ||||| ||||| ||||| ||||| |||||
XX 195 rGlySerArgAspArgSerArgAlaSerArgSerArgSerArgGlySer 211

```

```

336 .....CTTAAGCGCGGANGAAN 352
212 ProLeuGlyAlaThrAlaThrSerCysProAlaArgArgArgCysSe 228
353 TNNNGNCATTCATTCATTCGGTTGTGAGCTGGCGCTGACCCCT 402
228 rIleGlyAlaSerSerGly.....CysProHisProP 239
403 CGCTTACGCAAAATCCCTGC.....CGTCGATGC 431
239 rovalArgArgSerProvalAsnSerSerLysArgAlaHisArgArgCys 255
432 CGAGCCGTTGCC.....CATCT 448
256 ThrAlaArgArgGlyArgPheArgGlyProHisArgAspThrGlyArg 272
449 TCGTCGAATCGCATGAGCAGCAATCCGCTNGCGGAGACCTGGTGTG 498
272 gArgArgCys.....TrpArgTrpProArgProArgArgCysArgCys 287
499 ATCAAGAAGAGCGGANGANGATTCAGAGANGTGTCTGTATTTAGCCG 548
287 eArgArgTrpGlyArgProLeuTrpAla..... 296
549 TTTGACCGGCGTAAATCCATGCTGTAGAGCAGCTGGCCGACA..... 593
297 .....SerGlyCysProArgAlaArgTrpArgArgGlySe 308
594 .....CGTCCGCTGAAAATGCTGCC 615
308 rAsnTrpSerSerGlyArgSerSerAlaAlaSerProLysArgArgCysG 325
616 AACATGCA.....AACACATGAATTCGGCG.....CCGCA 647
325 LArgArgValArgSerAspThrSerAlaArgArgSerArgCysProAla 341
648 TCGGCGCGTTGAGTGCAGCAGCAGCATTCATTATGAGCGGCGGTG 697
342 SerSerProLleArgTrp.....ThiGlyArgCys 351
698 CAACAAAACCGTTTGGACCATCAATTATCA..AGATGTAATTGGCATC 744
351 sArgArgTrpArgArgProLeuGlyCysSerProAlaArgAlaHisCys... 366
745 GGAAGCTTTGTTGCAACAGCGCTCGAACACCGAGCGGTATGCTT 794
367 ..ThrAlaArgCysGlyArgAspGly.....CysSer 376
795 GGGGCTTCGACGATCAACAAACACGCGCTTGTGCTACCGTTTGGGTG 844
377 AlaPhePheGlyAsnProLeuHisArgSerLeu..... 387
845 CGAAAGTATGCAAAATTACTGCGGCGAATTGTTGACGACAGACCGC 894
388 .....ArgGlyProT 391
895 GTGATTCGCGTTCGATATGACGG...CGGATTCACAGCGCGCA 941
391 rPalaAlaProPheArgAlaHisArgSerArgSerThrThrArgArgCys 407
942 CGATTATTTGGAGCGCTACCAATAGATTTCCGTTATGAGAGAGCC 991
408 AlaValArgGlySerSerArgHisArgTrpAlaSerThrArgArgTrp 424
992 GCAGCAAGAGCTGTTGGCTGGTGGCC..... 1022
424 OHHisProPro.....LysGlyCysAlaThrAspPheHisSerGlyA 439
1023 .....GCAGCCGCAAAATCTCATCAGCGGTAC 1052
439 rGlyCysTrpProArgThrAlaSerSerArgAlaAlaSerGlyAlaSer 455
1053 GACCTCGGCAATTTCTGAAAACAAACTCTTAAGTTACGACGACGCC 1102

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456 AlaLysArgThrArgLeuArgArgSerCysProValArgSerProAr 472
1103 TCNAACG.....TGG..... 1112
472 gArgArgGlyThrArgAlaAlaTrpHisSerAlaCysGlySerSerSerA 489
1113 ..CGAGCGCGCATGCTCGCATTTGGTACTTACGAGCGCGTAATGCC... 1157
489 rArgArgProSerSerGlyArgProTrpSerValProLleArgProSerSer 505
1157 ..... 1157
506 IleCysGlyArgAlaValGlyLeuThrSerProSerSerProLeuAsnAr 522
1158 .....GCTGACAT.....CCTGCCTACCTGCTTTTGGCCGATTTA 1194
522 gProPheAlaArgSerAlaProAlaSerThrProCysArgArgHisA 539
1195 ATCGTCGCGCATACCGCAGCAGCGCAAGCATTGGGTTGCTTGGAAATTGA 1244
539 snArgArgArgTrp.....Gly 544
1245 CGAGAGAGACCT 1256
545 SerArgArgPro 548

seq_name: /SIS1/gcgdata/geneseq/geneseqp-emb1/AA1999.DAT:AA04998
seq_documentation_block:
ID AA04998 standard; Protein; 388 AA.
AC
XX
AC AAY04998;
XX
DT 06-JUL-1999 (first entry)
XX
DE Mycobacterium species protein sequence 50B.
XX
KW Secreted protein; Mycobacterium; primer; PCR; amplification; probe;
XX hybridisation; detection; vaccine; immunisation; infection.
XX
OS Mycobacterium sp.
XX
PD W09909186-A2.
XX
PD 25-FEB-1999.
XX
PE 14-AUG-1998; 98WO-FR01813.
XX
PR 11-SEP-1997; 97FR-0011325.
XX
PR 14-AUG-1997; 97FR-0010404.
XX
PA (INSP ) INST PASTEUR.
XX
PI Gicquel B, Llm EM, Pellicic V, Portnoi D, Goguet de la Salmoniere Y;
PI Guigueno A;
XX
DR WPI; 1999-181045/15.
XX
DR N-PSDB; AAX34249.
XX
PT Mycobacterial DNA vectors containing reporter constructs - for
PT identifying coding or promoter sequences involved in
PT infection-associated protein expression
XX
XX
XX Claim 32; Fig 50B; 309pp; French.
XX
XX Sequences AAY04742-Y05000 and AAY07201-Y07204 represent secreted
XX proteins from various Mycobacterium species microorganisms. The
XX encoding nucleotide sequences can be used as primers and probes for
XX methods for detecting and identifying mycobacteria, especially belonging
XX to the M. tuberculosis complex. The encoded proteins can be used in
XX vaccines for immunisation against a bacterial or viral infection.
XX
XX

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50 Sequence 388 AA;

alignment_scores:
Quality: 131.00 Length: 355
Ratio: 0.873 Gaps: 20
Percent Similarity: 42.254 Percent Identity: 27.042

Alignment block:

US-09-303-518D-127 x AAY04998

Align seg 1/1 to: AAY04998 from: 1 to: 388

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405 GTTCGCAAAATCCCTGCGTCGATGCCGAGCGGTCGCCATCTTCGTCGA 454
      ::::::::::::::::::::|::|::|::|::|::|::|::|::|::|::|
1 11eA9d9g1ygl1yCysArGArGArG1yTrpArGser...ArGAr 16
      ::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|
455 ATGCGATGAGACCAATCCGTCGCGGAGACCCCTGTGGTGTGATCAAA 504
      ::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|
16 gCysG1yArGcysArGArGAlaValG1y.....A 26
      ::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|
505 GAACCCGCGANGATTTTCAGACGANGTTCGTATGAGCGGTTTGAC 554
      ::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|
26 rGArGArGArGArGArGTrpG1n..... 32
      ::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|
555 CGAGCGTAAATCCATGTGTGTAGGACGACGCGCAGACGTCGCGTCTG 604
      ::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|
33 .....TTrpArG1uArG..... 36
      ::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|
605 AAATGCTGCGCAACATCGAAACATGAATTCGGCGCGCGCGATCCGGCG 654
      ::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|
37 ArGArGcysG1n1sArG1nArGTrpArGArGTrpArGArGcysA 53
      ::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|
655 GGTTCGTCGCGCGCGACATTCATTTCATGACGCGGTCGTCGAACAA 704
      ::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|
53 rGTrpArGTrpArGmet.....Ala1eArGArGArGArG 64
      ::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|
705 AACCGTTTGGACCATTCATTAAGATGTATTCATCGGACGTTTGT 754
      ::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|
65 G1yArGArGTrhArGArGTrhArGArG..... 73
      ::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|
755 TTGCAACAGCGCGCTGTAACACGCGCGT..... 785
      ::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|
74 ....ASnArGPrArGArGArGArGArGArGArG1yPrArG1yArG 89
      ::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|
786 .....GATTCGTTGGTGTCTCAAGTCGAACAAACACG..... 821
      ::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|
89 rGArGPrArGTrhArG1yG1yTrpArGArGArGTrhArGArGAla 105
      ::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|
822 .....CTTCCTGCGTACCGTTTGGTGGTCGGAAGTAT 853
      ::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|
106 ArGArGTrpTrpArG1yPrArGAla1nArGserG1yArG1n1s.... 120
      ::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|
854 CGCAATATACGCGCGCAATTCGTCGACGACGAACCGCGTATTC 903
      ::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|
121 .....G1yArGArG1yTrpArGArGTrpAlaArGArG1n 133
      ::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|
904 GGTTCGATTCGAACGCGCG.....GATTAC 929
      ::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|
133 rGArG1yArGArGArGArGArGcysArGArGAlaArGArG1yArGArG 149
      ::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|
930 ACAAGCGCGCGACGATTAATTCGACGCGTACCAATCATGATTCGTTA 979
      ::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|
150 SerArGArGcysArGArG1uArG1n1sAlaArGArGAlaArGArGAr 166
      ::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|
980 TCGAAGAAGCGCGCGACA.....AGACGTTCGCGCTG 1014
      ::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|
166 gArGArGArGArGArG1nG1nPrArG***TrpArG1yArGArGArG 183
      ::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|
1015 GT...TTCGCGCGACGCGCAATATCTCCATCCAGCGCGACGCGCTCG 1061
      ::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|
183 1yArGcysArGArGAlaPrArG1n1sArG1nTrpArGAlaArGArGPrArG 199
      ::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|

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1062 CCATTCTCTGAAAAACAACCTTCAGTTCACGACGCGTCACGGTG 1111
      ::::::::::::::|::|::|::|::|::|::|::|::|::|::|
200 .....ArgSerArG1n1sG1yArG1n1sArG1yArGTr 211
      ::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|
1112 GCGACCGCGC.....CATGTCGCCAT.....TGGTACT 1140
      ::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|
211 pArGArGArGArGArGTrp1yArG1n1sAlaArGArGPrArGArG 228
      ::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|
1141 TACGAGCGCGTAATCCGCTAGACATCT...GCCATCCCTGCT...TTT 1184
      ::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|
228 rGArGcysArGArG1yPrArGArGser1eueG1yTrpPro***ArGAr 244
      ::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|
1185 GCGGATTTATGTCGCGCGATAC.....CGACGCGCGCAAG 1222
      ::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|
245 AlaArGTrp1uArG1yArGTrpArGPrArGAla1eArG1nArGArG... 260
      ::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|
1223 CATGCGGTCCTTGATTCAGTAAGAACCTCCCTTTGCGACGCTTC 1272
      ::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|
261 .....ArGArGArGPrArG1n1sArGArGAsNT 270
      ::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|
1273 GTTCGCGCGCGCAATACGAATANGCCCGCTGTCGTAAGTCGTCGA 1322
      ::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|
270 hrAlaG1yG1yG1uArG1l1eG1yAspG1yPhe..ValArGcysTrhA 286
      ::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|
1323 AACCNFTGAGAAAG 1336
      ::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|
286 rGPrArGTrhArGArG 290
      ::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|

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seq_name: /SIDS1/scgdata/geneseq/geneseq-emb1/AA2001.DAT:AA59827

seq_documentation_block:

ID AA59827 standard; Protein: 1592 AA.

AA59827;

04-APR-2001 (first entry)

Protein #4 encoded by Tuid/E gene.

Toluene degradation; enzyme: waste degradation; Tude; Tuid.

Thauera aromatica.
Xanthomonas maltophilia.
Geobacter metallireducens.
Azotarcus toluilyticus.

W0200072650-A2.

07-DEC-2000.

24-MAY-2000; 2000MO-US14298.

01-JUN-1999; 99US-0323872.

(UYOH-) UNIV OHIO.

Coschigano PW;

WPI: 2001-041080/05.

N-PADB; AAF23627.

Composition comprising toluene degrading enzyme useful for biological treatment of organic compounds, especially for degrading toluene or its analogs

Disclosure: Fig 12; 122pp; English.

The present invention relates to toluene degrading enzyme genes and proteins tuid (see AAF23629 and AAB59831), tui1 (AAF23630 and AAB59832), tuf (AAF23631 and AAB59833) and tucG (AAF23632 and AAB59834). The toluene degrading enzymes are homologues of pyruvate formate lyase. The toluene degrading enzymes are useful for biological treatment of organic compounds and in particular for the degradation of toluene and its


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875 ..AATTGCC... ..GCAGTAATTGCGATTCTTGCACCCCAA 837
      |||||
252 aGlnSerProLeuAlaThrAlaIaSerAlaSerThrSerAlaProVal 268
      |||||
836 ACGGACGACAGAGCGCGTGTGTGACTTGAGACACCACCAAGCAAT 787
      |||||
269 SerCysGlySerSerAlaSerLeuAlaGlyProHisProGlyThrSe 285
      |||||
786 CACGGCTCGGTGTTCAGACGCGCTGTGCAACCAACCGTCGATGGCAA 737
      |||||
285 rAspLeuHisIleSerSerThrProAlaIaIaThrThrLeuPro..... 299
      |||||
736 TTAACATCTGATTAATTGATGTCACAAAGGTTTGTTCACGACGCGC 687
      |||||
300 .....ValMetIleGlyThrGluProThrSerProThrPro 311
      |||||
686 TCAATGAATGATGTCGTCGACACACAA...CCGCGCGGATGCGCGGC 640
      |||||
312 Ser.....AlaPheLysGlyProSerHisSerGlyAs 322
      |||||
639 GCGCAAT.....TCATGTGTTGATGTCGACGACATTTTCAG 602
      |||||
322 nProSerHisGlyThrLeuGlyLeuSerGlyThrLeuGlyAlaTyrT 339
      |||||
601 ACGGACGTCGCGCGCGACGCTGCTACACACATGATTTTACGCTCGGTC 552
      |||||
339 hSerThrSerValProIleSerLeuSerAlaCys..... 350
      |||||
551 AAACGGCTCAATACACAGCANACNTCTGAATCTGCGCGCTTCTT 502
      |||||
351 .....LeuAsnProAlaLeuSerGlyLeuSerSerSerThrProle 365
      |||||
501 GATCACACACACAGGCTGCGCGACGCGATGTCATGCGATGCA 452
      |||||
365 u..... 365
      |||||
451 CGAAGATGGGACAGCGCTCGCATCGACGACGAGGATTTTGTGACGGA 402
      |||||
366 .....AsnGlySerAsnProLeuSerSerIleSerLeuProPro 378
      |||||
401 CGGNTACGACGCGACGATCCAAACCGGATGATCAGATTGCGCNCNNAN 352
      |||||
379 HisGlySerSerThrProIleAlaProValPheThrAlaLeuProSerPh 395
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351 TTGNTGCGCGCTTAAGTTGCCAAGCTTCGCGCGCGCTTGCAGACT 302
      |||||
395 eThrSerLeuThrAsnAsnPhProLeuThrGlyAsnProSerLeuAsnP 412
      |||||
301 CGATTTGCTGCTTGCCTTCAACGCGCAATACGACGATGAATGACGCGC 252
      |||||
412 rSerValSerLeuProGlySerLeuIleAlaThrSerSerThrAlaAla 428
      |||||
251 TTTTCG.....CCGCGATGATGCGCGGCGGATTTTG... 222
      |||||
429 ThrSerThrSerLeuProHisProSerSerThrAlaAlaValLeuSerG 445
      |||||
221 .....CCTGAACNGGGCG..... 207
      |||||
445 yLeuSerAlaSerAlaProValSerAlaAlaProPheProLeuAsnLeuS 462
      |||||
206 .....GTAAACACACGCGCGGATGCTTGTGCT 177
      |||||
462 eThrAlaValProSerLeuPheSerValThrGlnGlyProLeuSerSer 476
      |||||
176 TCAACACGACTTGCGCTTTTTCAGCGCATCG.....CCTTCCTTGAC 133
      |||||
479 SerAsnLeuSerTyrProGlyPheSerValSerAsnThrProSerValTh 495
      |||||
132 TTTCAATCNAAGGCGCATACCGCATATCTTCGCAAGACGCGGCTT 83
      |||||
495 rProAlaLeuProSerPheProGlyLeuGlnAlaIaProSerThrValAla 512
      |||||
82 CGGTA.....ATGACGGGCGCCGCTCA 63

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      |||||
512 IaValThrProLeuProValAlaAlaIaThrAlaProSer 524

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seq_name: /SIDSI/gcgdata/geneseq/geneseq.embl/AA2001.DAT.ABG03731

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seq_documentation_block:

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ID ABG03731 standard; Protein; 696 AA.

```

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AC ABG03731;

```

```

DT 13-FEB-2002 (first entry)

```

```

DE Novel human diagnostic protein #3722.

```

```

KW Human; chromosome mapping; gene mapping; gene therapy; forensic;

```

```

KW food supplement; medical imaging; diagnostic; genetic disorder.

```

```

OS Homo sapiens.

```

```

PN W0200175067-A2.

```

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PD 11-OCT-2001.

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PF 30-MAR-2001; 2001WO-US08631.

```

```

PR 31-MAR-2000; 2000US-0540217.

```

```

PR 23-AUG-2000; 2000US-0649167.

```

```

FA (HSE-) HXSEO INC.

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```

PI Dyrmanac RT, Liu C, Tang YT;

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DR WPI: 2001-639362/73.

```

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DR N-PSDB; AAS67918.

```

```

PT New isolated polynucleotide and encoded polypeptides, useful in

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PT diagnostics, forensics, gene mapping, identification of mutations

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PT responsible for genetic disorders or other traits and to assess

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PT biodiversity

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PS Claim 20; SEQ ID No 34090; 103bp; English.

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CC The invention relates to isolated polynucleotide (I) and

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CC polypeptide (II) sequences. (I) is useful as hybridisation probes,

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CC polymerase chain reaction (PCR) primers, oligomers, and for chromosome

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CC and gene mapping, and in recombinant production of (II). The

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CC polynucleotides are also used in diagnostics as expressed sequence tags

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CC for identifying expressed genes. (I) is useful in gene therapy techniques

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CC to restore normal activity of (II) or to treat disease states involving

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CC quantitating a polypeptide in tissue, as molecular weight markers and as

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CC a food supplement. (II) and its binding partners are useful in medical

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CC imaging of sites expressing (II). (I) and (II) are useful for treating

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CC disorders involving aberrant protein expression or biological activity.

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CC The polypeptide and polynucleotide sequences have applications in

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CC diagnostics, forensics, gene mapping, identification of mutations

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CC responsible for genetic disorders or other traits to assess biodiversity

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```

CC and to produce other types of data and products dependent on DNA and

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CC amino acid sequences. ABG00010-ABG30377 represent novel human

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CC Note: The sequence data for this patent did not appear in the printed

```

```

CC specification, but was obtained in electronic format directly from WIPO

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CC at ftp.wipo.int/pub/published_pct_sequences.

```

```

XX Sequence 696 AA;

```

```

alignment_scores:

```

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Quality: 121.00

```

```

Ratio: 0.571

```

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Percent Similarity: 45.396

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Length: 467

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Gaps: 23

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Percent Identity: 23.555

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alignment_block:

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US-09-303-518D-127 x ABG03731 ..

Align seg 1/1 to: ABG03731 from: 1 to: 696

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9 AATGAAAGGCTGTAACCTGACCATCGGCGAGACCGAGCACTCA 58
14 AsnLeuArgLysProSerLysArgLysAspArg***ArgAr 30
59 TTATGACGGCCCGTCATTACCAAGTCGCGTTCGGCGAAGATAT 108
30 gValSerProThrArgSerGlyLysArgArgGlyAla..... 42
109 GCCGGTATGCGCCCTNGATGAAGTCAAGGACCGATGCCGTAATA 158
43 .....GluGlyLysAsnArgGlnGlyLysLysLys 52
159 AGCCCAAGTGTGTTGAAGACAAAGAAAGATCCGGCGTGTTCACG 208
53 GluArgGlyLysGluArgArgGlyLysArgSerGluArgGlnArgAspAr 69
209 CGCCAGTTTCAGGCAAAATCCGCGCATCCATCCGCGGCAAGACGCGTA 258
69 gArgArgArgLysGlnArgLysGlnGluGlnArgArgArgAlaArgT 86
259 CTTCAGTCGTCGTGATTGCCGTTGAAGCAACGCAAGATCGAGTTGCA 308
86 hAsn.....GluArgLysProArg..... 92
309 AGCTACGCGCGGAGAGCGTTGGCAACTTAAAGCGCGCAGNANTNGNN 358
93 .....GlnThrGlnAlaAsnGlyAlaThrSerSer***LysAlaSerAl 107
359 GCAATCTGATCCATCCGCTTGTGTGACTGCGCTGCGTANCGTCGCTGC 408
107 aGlnGlnAlaGlyMet.....TyrGlyLysSerPro***IleA 120-
409 AGCAAAATCCCTGCGTGCATCGGAGCGCGTCCGCAATCTTCGTCAATGC 458
120 sPAlaThrAlaIleArgArgGlyGlyAlaProCysSerArgArgThr 136
459 GATGACACCAATCCGCTNGCGGAGACGCTGTGTTGATCAAGAAG 508
137 CysLeuAsnGlnGlyThrIleAlaThrProSerGly.....Ar 149
509 CGGNCAGNATTTCAAGACGANGTTCGTATGAGCCGTTGACCGAG 558
149 gArgArgGlnGlyAspAlaGly***ProGlyLysLeuAlaSerGlnHisAsp 165
559 CGTAAATCCATGCTGTGAAGCGAGCGAGCGAGACGTCGCTGGAAGA 608
166 AlaSerGlnHisGlyCysLeuArgThrGlnAlaGly***ProSerAsp 182
609 T.....GCTGCAACATCGAAGACATG 631
182 rThrGlnSerValCysArgArgProLeuAlaMetHisValProThrHisG 199
632 AATTGCGCGCGCGCATCCGCGGTTGATGAGCGAGCAATC..... 676
199 lSerHisGlyProValPheThrArgLeuValSerHisThrPheHisCys 215
677 .....ATTTCATTGACGCGTGTGCAAAACAAACCG...T 710
215 sGlySerLysLeuProAlaValGlyArgProValAlaCysArgProThrT 232
711 TTGACCATCAATATTCAGATGTAATTTGCCATCGACGTTGTTGCA 760
232 ySerProSerLeu.....CysHisAsnPro..... 81
761 CAGGCGCTGTGAACCGAGCGGTGATGCTTGGGTGTTCTCAAGC 810
241 nArgProAlaGlnLeuLeuAlaHisSer.....SerAla 253
811 AACAAACCAAGCCTTTCGTACCGTTTGGTGCAGAAAGTATCGCAAT 860

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253 euGlnCysAlaProLeuSerTyrAsp.....ProGln 263
861 TACTGCGGCGCAATTGCTTGACGAGACAAACCGGATTTCCGGTGG 910
264 ArgCysAla.....ProProSerProArgProHisArgArgL 276
911 TATTGAACGCGCGGATTACCAAGCGCGCAGATTAATTTGGAGCGTAC 960
276 yProProSerProHisProHisArgArgAla.....P 287
961 CACATTCAGATTCCGTATGCA..... 983
287 rProSerProHisProHisArgArgAlaHisThrThrAlaArgThrAsp 303
984 .....AGAGGCGCGCAGCAAGACGCTGTCG 1009
304 ProThrThrSerAlaProProProAlaGlnThrGlnArgArgAlaThrAr 320
1010 GCTGGGTTGGCGCGCAGCGCAAAATCTCATCGCGGTACGACCTTC 1059
320 GluProAlaThrLysHisThrArgAsnAlaHis.....ProA 333
1060 GCCCATTTCTGAAAGCAAACTCTTCAGTTCACGACAGCGCTGCAAG 1109
333 rArgSerAlaCysAsnArgGlyThrHisThrHisProArgArgArgArg 349
1110 TGG.....CGACCGCGCAT.....GATGC 1129
350 ThrThrGluArgThrThrHisHisAlaArgProArgAsnArgLysGlnAl 366
1130 CGATTGTGTTACTTACGAGCGGTAATGCCGCTAGACATCTGCTACCCCTG 1179
366 aThrProAsnThrArgGlnProThrAlaGlyArgHisGlyGluThrAspG 383
1180 CTTTGGCGCGATTAACTCGTCGGGATACGACAGCGCGCAACATTTGG 1229
383 lAlaThrArg.....ArgArgGlnHisGlyGlnThrArgGlyGly 397
1230 TTGCTTGGAATTGGAGCAGAGAACGCTGCTTGTGCGACCTGCT 1274
398 .....GlyArgArgArgGlyArgAlaAlaLysThrArg 408

seq_name: /SIDSI/gcgdata/geneseq/geneseqp-emb1/AA2001.DAT:ABG23389
seq_documentation_block:
ID ABG23389 standard; Protein; 1194 AA.
XX
AC ABG23389;
XX
DT 18-FEB-2002 (first entry)
XX
DE Novel human diagnostic protein #23380.
XX
KW Human; chromosome mapping; gene mapping; gene therapy; forensic;
KW food supplement; medical imaging; diagnostic; genetic disorder.
XX
OS Homo sapiens.
XX
PN W0200175067-A2.
XX
PD 11-OCT-2001.
XX
PE 30-MAR-2001; 2001WO-US08631.
XX
PR 31-MAR-2000; 2000US-0540217.
PR 23-AUG-2000; 2000US-0649167.
XX
PA (HYSE-) HYSEQ INC.
XX
PI Drmanac RT, Liu C, Tang YF;
XX
WP1; 2001-639362/73.

```

DR N-PSDB; AAS87576.

XX New isolated polynucleotide and encoded polypeptides, useful in
PT diagnostics, forensics, gene mapping, identification of mutations
PT responsible for genetic disorders or other traits and to assess
XX biodiversity

PS Claim 20: SEQ ID No 53748; 103pp; English.

XX The invention relates to isolated polynucleotide (I) and
CC polypeptide (II) sequences. (I) is useful as hybridisation probes,
CC polymerase chain reaction (PCR) primers, oligomers, and for chromosome
CC and gene mapping, and in recombinant production of (II). The
CC polynucleotides are also used in diagnostics as expressed sequence tags
CC for identifying expressed genes. (I) is useful in gene therapy techniques
CC to restore normal activity of (II) or to treat disease states involving
CC quantitating a polypeptide in tissue, as molecular weight markers and as
CC a food supplement. (II) and its binding partners are useful in medical
CC imaging of sites expressing (II). (I) and (II) are useful for treating
CC disorders involving aberrant protein expression or biological activity.
CC The polypeptide and polynucleotide sequences have applications in
CC diagnostics, forensics, gene mapping, identification of mutations
CC and to produce other types of data and products dependent on DNA and
CC amino acid sequences. ABG00010-ABG30377 represent novel human
CC diagnostic amino acid sequences. ABG00010-ABG30377 represent novel human
CC Note: The sequence data for this patent did not appear in the printed
CC specification, but was obtained in electronic format directly from WIPRO
at ftp.wipro.int/pub/published_pct_sequences.

SO Sequence 1194 AA:

alignment_scores: Quality: 113.50 Length: 441
 Ratio: 0.565 Gaps: 18
Percent Similarity: 45.578 Percent Identity: 21.995

alignment block:

US-09-303-518d-127/rev x ABG23389

Align seg 1/1 to: ABG23389 from: 1 to: 1194

1313 TTACGACACGAGGGGCGCCATTCGTTTGGCCGGGACACGACGACTGCA 1264
||| :|||:|||||
584 LeuSerGlnSerGlyProProGlyLeuLeuPro..... 594
1263 CAAGGAGAGGTCTTCTTCGTCATTCACGACCAACCATGCT...TGCG 1217
||| ||| :|||:|||||
595SerProSerPheAspSerIysProProThrThrLeuLeuG 608
:|||| :|||:|||||
1216 CGGTGCGGTATCGCGGACGATTAAATCGCGCAAAAGCAGGGTAGCGAGG 1167
:|||| :|||:|||||
608 LysLeuLeuProAlaProSerMet..... 615
1166 AGTGTACGCGGATTACGCGCTGTAAGTACCAATCGGACCATGGCGG 1117
||| ||| :|||:|||||
616ValPro...AlaThrAspThrLy 622
1116 GTGCGCACCGTTGACGCGTGTGTAAGTGAAGATTGTTTTCACAGA 1067
:||||:|||||
622 sAlaProProThrLeuGlnAlaGluThrAlaThrIysProGlnAlaThrS 639
1066 AATGGCCGAGGCGTACGCGGTAGAGATTTTCCGGCGCGCGCA 1017
:|||| :|||:|||||
639 eAlaIleProSerProAlaProIysGlnSerPheLeuPheGlyThrClnAsn 655
:|||| :|||:|||||
1016 ACCGACCGACAGCTCTTGTGCGGCTCTTGATGATACGGAATGTG 967
:||||:|||||
656 ThrSerProSerSer.....ProAlaAla..... 663
966 ATTGTGTAGCGCTCCCAATATATCGTGGCGCTTGTGTATCGCGCGT 917

664ProAlaIleSerSerAlaProProMetPheIysProI 676
:||||:|||||
916 TCATATCCGACCGGAAATACGCGGTGTGTGCGCAACCAATTCGCCG 867
:||||:|||||
676 IePheThrAlaProProIysSerGlnIysGlnGlyProThr..... 689
866 GCAGTATATTTGGCATATCTTCCGCCACCAAAACGGTAGCGAAGAGCGTGG 817
689 689
816 TTGTGTAAGTGAAGACCCCAAGCAATCAGCGGTGCGGTTCACAC 767
:||||:|||||
690ProProGlyProSerValThrAlaThrAla..... 699
766 GCGCTGTGCAAAACAGCGTCGATGCGAATTCATCTTGATTAATGATG 717
||| :|||:|||||
700 ..ProSerSerSerSerLeuPro..... 706
716 GTCCAAACGGTTTGTGTCACCGCGGCTCATGAATGAATGTGCGT 667
:||||:|||||
707 ThrThrThrSerThrThrAlaProThr..... 715
666 GCCACTCAACCG.....GCCGATGCGGCGCGCGCAATTCATGTGTT 623
:||||:|||||
716PheGlnProValPheSerSerMetGlyProProAlaSer...ValP 730
622 CGATGTGCGACATTTTTCACAGCGGACGTCGTGCGCAGCTGCTTACAC 573
:||||:|||||
730 IeLeuProAlaProPhePheIysGlnThrThrProAlaThrAlaPro 746
572 ACATGATTTTACGCTCGTCAACGCGCTCAATACGACCAATCTGCT 523
|||
747 Thr..... 747
522 GAATCTGCGCGCTTCTTGTATCACAACACGAGGCTGCGCGNAGCG 473
:||||:|||||
748ThrThrAlaProLeuPheThrGlyLeuAlaSerAlaThrSerA 762
472 GATTGTCGATCGCATTTGACGAGGATGCGGACGCGCTGGCGATCG... 426
:||||:|||||
762 IeValAlaIleProIleThrSerAlaSerProSerThrIlePheSerLys 778
425 ..ACGGAGGAGATTTGCTGTAACGAGCGGTAACGAGCGCAGCCACAA 379
:||||:|||||
779 ProAlaPheGlyPheGlyIleAsnSerValSerSerSerSerVal...Se 794
378 ACCGATTTGATCGATTTGCTGACGCGCNCNNANTGCGCGCTTAAGTTGCC. 330
:||||:|||||
794 ThrThrThrThrSerThrAlaThrAlaIleSerGlnProPheLeuPheGlyA 811
329AACGCTTGGGCGCGTAGCGGTGCAAGTCGATTGCTGCTG 288
:||||:|||||
811 IeProGlnAlaIleSerAlaIleSerPheThrProAlaIleGlySerIlePhe 827
287CCTTCAACGGCAATCAGACCGAGTGAAGTACGCGCTT 250
:||||:|||||
828 GlnPheGlyIysProProAlaIleProThrThrThrThrValThrThrPh 844
249 TTGCGCCGCGATGATGCGCGGATTTTGCCTGAACGCGCGGTAAACA 200
:||||:|||||
844 eSerGlnSerIleuHisThrAlaValProThrAlaThrSerSerSerAlaA 861
199 CCAAGCGCGGATCTTGTGCTTCAACAGCACT...TGCGCTTTTGG 153
:||||:|||||
861 IeAspPheSerGlyPheGlySerThrLeuAlaThrSerAlaProAlaThr 877
152 ACGGATCGCTTCTGCTGATTCATCAGGCGCGCATACCGGACATATTC 103
:||||:|||||
878 SerSerGlnProThrLeuThrPheSerAsnThrSerThrProThrPheAs 894
102 TTGCGCAGACGCGACCTTCGTTATGACGGCGCGCTATATAATGACTT 53
:||||:|||||

894 nileProheGlySerSerAlaIysSerProLeuProSerTyrProGlyA 911
 52 GCTCCGGTCTGCGCGCATGAGC 30
 : : : : :
 911 laasnProGlnProAlaPhedGly 918

seq_name: /sids1/gcgdata/geneseq/geneseq-emb1/AA2001.DAT:AAU39645

seq_documentation_block:

ID AAU39645 standard; Protein; 338 AA.

AAU39645;

13-FEB-2002 (first entry)

Proionbacterium acnes immunogenic protein #541.

XX SAPHO syndrome; synovitis; acne; pustulosis; hypertosis; osteomyelitis;
 KW uveitis; endophthalmitis; bone; joint; central nervous system; ELISA;
 KW inflammatory lesion; acne vulgaris; enzyme linked immunosorbent assay;
 KW dermatological; osteopathic; neuroprotectant.

XX Proionbacterium acnes.

MO200181581-A2.

01-NOV-2001.

20-APR-2001; 2001MO-US12865.

21-APR-2000; 2000US-199047P.

02-JUN-2000; 2000US-208841P.

07-JUL-2000; 2000US-216747P.

(CORI-) CORIXA CORP.

Skeiky YAM, Persing DH, Mitcham JL, Wang SS, Bhatia A;
 L'maisonneuve J, Zhang Y, Jen S, Carter D;

WPI; 2001-616774/71.

N-PSDB; AAS59508.

Example 1: SEQ ID NO 840; 1069pp; English.

Sequences AAU39105-AAU68017 represent Proionbacterium acnes immunogenic

polypeptides. The proteins and their associated DNA sequences are used in

the treatment, prevention and diagnosis of medical conditions caused by

P. acnes. The disorders include SAPHO syndrome (synovitis, acne,

pustulosis, hypertosis and osteomyelitis), uveitis and endophthalmitis.

P. acnes is also involved in infections of bone, joints and the central

nervous system, however it is particularly involved in the inflammatory

lesions associated with acne vulgaris. A method for detecting the

presence or absence of P. acnes in a patient comprises contacting a

sample with a binding agent that binds to the proteins of the invention

and determining the amount of bound protein in the sample. The

polypeptides may be used as antigens in the production of antibodies

specific for P. acnes proteins. These antibodies can be used to

downregulate expression and activity of P. acnes polypeptides and

therefore treat P. acnes infections. The antibodies may also be used as

diagnostic agents for determining P. acnes presence, for example, by

enzyme linked immunosorbent assay (ELISA)

Note: The sequence data for this patent did not form part of the printed

specification, but was obtained in electronic format directly from WIPO

at ftp.wipo.int/pub/published_pcl_sequences.

Sequence 338 AA:

alignment_scores:

Quality: 112.00 Length: 375
 Ratio: 0.747 Gaps: 18
 Percent Similarity: 40.000 Percent Identity: 22.667

alignment_block:

us-09-303-518D-127/rev x AAU39645

Align seg 1/1 to: AAU39645 from: 1 to: 338

1031 TCCGGCTGCGCGCAACCGCAACCGCAACGCTCTTCTGCGCGCTCTTC 982
 : : : : :
 17 ThrGlyCysAlaGlyLeuThrSerLeuSerSerThrThrArgAlaSerH 33
 981 GATAACGGAAATCTGATTGTGTAGCTGCCAAATAATATCGCGCCCTT 932
 33 Ser : : : : : SerCysValProV 39
 931 GTGTAATCGCGCGCTTCATATACGACCGGAATATACGCGGTGTCTGCG 882
 : : : : :
 39 Al : : : : : ThrSerProGlySerSerSerSerSerSer 49
 881 TCAACCAATTCGCGCGCAATTTGGATATTTGGATATCTTCGCCCAAAACG 832
 : : : : :
 50 ProThrThrThrSerAlaAlaAlaArgArgArgLeuArgProArgSerP 66
 831 ACGCAAG : : : : : AGCGGTGTTTGTGATCTGAG 803
 : : : : :
 66 OThrLeuThrThrCysSerGlyTTProArgArgSerLeuProAlaAlaP 83
 802 AACCAACCAACCAATTCACG : : : : : CGCTCG 777
 : : : : :
 83 roAlProThrSerProThrThrArgAlaArgAsnThrProProGlnSer 99
 776 GGTTCACAGCGCGCTGTGCAACCAACCGTCCGATGCAATTCATCTTG 727
 : : : : :
 100 AlaThrArgArgPheProThrSerLeuThrProLeuArgThrThrSer 115
 : : : : :
 726 ATAAATGATGTCACAAACGCTTTGTTGACCGCAGCGCTCAATGAAT 677
 : : : : :
 116 : : : : : ArgCysLeuAlaThrAlaArgProGly 124
 676 GAATGCGCTGCCACTCAACACGCGCGGATG : : : : : GGCGCGCGAAT 633
 : : : : :
 125 : : : : : TyrSerLeuSerThrThrSerSerThrCysArgValGlyProSerSer 140
 : : : : :
 632 TCATGTGTTTCGATGTGCA : : : : : 612
 : : : : :
 141 AlaCysValThrAlaLeuSerAlaGlyIleThrValThrThrArgSerG 157
 : : : : :
 611 : : : : : GCATTTTCAGACGCGCTGCGCGACCTGCCCTTA : : : : : CACACAT 569
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 157 uCysAlaLeuThrProSerThrSerProProAlaThrGlyPheThrT 174
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 568 GGATTTTACCGCTGCTCAACAGGCTCAATACGACGACGACGCTGTGAA 519
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 518 TCNTCGCGCGCTCTTGTATACACACGAGGTCTCCGACGCGAAT 469
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 175 : : : : : SerThrThrMetLeuThrThrSerGlyThr : : : : : 184
 : : : : :
 468 GGTTCATCGCATTCAGCAAGATGCGAAGGCTTCGGCATGACGAGGAG 419
 : : : : :
 185 : : : : : SerArgAlaLeuThrAlaIleArgArgLeuThrIleSerAlaAla 200
 : : : : :
 418 GGATTTGCTGAACGAGCGGNTACG : : : : : 393
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 200 rGlyThrProValValGlySerArgArgPheProArgProThrArgThrPro 216
 : : : : :
 392 : : : : : ACGCAGTCACCAACCGGATTTGATGAC 364
 : : : : :
 217 serCysGlyAlaMetProThrLeuArgIleSerArgProThrThrPmet 232


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363 ATTCGNCNNANTTCNTCGCGCTTAAGTTGCCAACGCTTGCGGCGGT 314
233 .....ProSerMetThrSerCysIleclYthrThra 243
313 AGCGTCGAACTCGATTTCGTGCTTGCCTTCAACGCGCATACGACGAC 264
243 rGArgGlySerProIleProSerLeu.....ThrAlaSerThrIleVal 257
263 TGAAGTACGCGCTTTGCGCGCGATGATG.....GGCGGAT 226
258 ArgSerSerArgArgSerValArgIrrPalArgSerSerSerSer 274
225 TTGGCTGAACNGCGCGGTAACACCGCCGATMC.....TTT 182
274 oHisProArgThrSerProThrGlyThrIleGlySerThrArgPhea 291
181 TGCTTCGAACAGCAGCTTGCGCTTTTGGAGCGATGCGCTTCCTGAC 132
291 rgProSerAlaArgSerTrp..... 297
131 TTGATCAGNGGCGCGCATACGCGCATTCCTGCGCAACGCGACTTC 82
298 .....SerSerProSerSerAlaThrTrh 305
81 G.....GTAATGACGGGC 69
305 rSerArgAsnGlyValMetThrGly 313

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seq.name: /SIDSI/gcgdata/geneseq-emb1/AA1999.DAT:AAV31745

seq_documentation_block:

ID AAV31745 standard; Protein; 430 AA.

XX AAV31745;

DT 22-NOV-1999 (first entry)

DE Mycobacterium tuberculosis specific DNA-encoded polypeptide.

KW Tuberculosis; infection; diagnosis; DNA probe.

XX Mycobacterium tuberculosis.

OS Key Location/Qualifiers

FT Misc-difference 4 /note= "encoded by TGA"

FT Misc-difference 6 /note= "encoded by TGA"

FT Misc-difference 20 /note= "encoded by TGA"

FT Misc-difference 29 /note= "encoded by TGA"

FT Misc-difference 54 /note= "encoded by TGA"

FT Misc-difference 64 /note= "encoded by TGA"

FT Misc-difference 69 /note= "encoded by TGA"

FT Misc-difference 89 /note= "encoded by TGA"

FT Misc-difference 99 /note= "encoded by TGA"

FT Misc-difference 114 /note= "encoded by TGA"

FT Misc-difference 119 /note= "encoded by TGA"

FT Misc-difference 129 /note= "encoded by TGA"

FT Misc-difference 159 /note= "encoded by TGA"

FT Misc-difference 169 /note= "encoded by TGA"

FT Misc-difference 182 /note= "encoded by TGA"

FT Misc-difference 182

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FT /note= "encoded by TGA"
FT Misc-difference 185
FT /note= "encoded by TGA"
FT Misc-difference 219
FT /note= "encoded by TGA"
FT Misc-difference 259
FT /note= "encoded by TGA"
FT Misc-difference 269
FT /note= "encoded by TGA"
FT Misc-difference 291
FT /note= "encoded by TGA"
FT Misc-difference 323
FT /note= "encoded by TGA"
FT Misc-difference 339
FT /note= "encoded by TGA"
FT Misc-difference 349
FT /note= "encoded by TGA"
FT Misc-difference 356
FT /note= "encoded by TGA"
FT Misc-difference 366
FT /note= "encoded by TGA"
FT Misc-difference 410
FT /note= "encoded by TGA"
FT Misc-difference 421
FT /note= "encoded by TGA"
FT /note= "encoded by TGA"

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EP945462-A1.

29-SEP-1999.

25-MAR-1998; 98EP-0302287.

25-MAR-1998; 98EP-0302287.

(COPL) CSIR COUNCIL SCI IND RES.

Kumar D, Srivastava BS, Srivastava R;

WPI: 1999-530042/45.

N-PSDB; AAX87940.

New nucleic acid molecules, useful for detecting and identifying

Mycobacterium tuberculosis

Disclosure; Page 15-27; 42pp; English.

The present sequence represents an amino acid sequence deduced from the Mycobacterium tuberculosis specific DNA fragment provided in AAX87940. This DNA fragment comprises a StuI-StuI fragment of M. tuberculosis genomic DNA and contains an insertion sequence-like element and repetitive sequences. The DNA fragment is useful as a probe, especially for detecting or identifying M. tuberculosis in clinical isolates and body fluids e.g. sputum, cerebrospinal fluid, pleural fluid, urine, gastric lavage, bronchial lavage, pericardial or lymph node aspirate (all claimed). It is also useful for restriction fragment length polymorphism analysis of M. tuberculosis isolates (claimed). The probe provides rapid and specific diagnosis of tuberculosis and M. tuberculosis infection.

Sequence 430 AA;

alignment_scores:

Quality: 110.00 Length: 446

Ratio: 0.625 Gaps: 23

Percent Similarity: 39.462 Percent Identity: 23.767

alignment_block:

US-09-303-518D-127 x AAV31745 ..

Align seg 1/1 to: AAV31745 from: 1 to: 430

44 GACCGAGCAAGTCATTATGACGGCGCGTCATTA.....CCGAA 84

DE Novel human diagnostic protein #4805.

XX Human; chromosome mapping; gene mapping; gene therapy; forensic;
KW Food supplement; medical imaging; diagnostic; genetic disorder.

XX Homo sapiens.

PN WO200175067-A2.

PD 11-OCT-2001.

PF 30-MAR-2001; 2001WO-US08631.

PR 31-MAR-2000; 2000US-0540217.

PR 23-AUG-2000; 2000US-0649167.

PA (HYSE-) HYSEQ INC.

PI Drmenac RT, Liu C, Tang YT;

DR WPI; 2001-639362/73.

DR N-PSDB; AAS69001.

PT New isolated polynucleotide and encoded polypeptides, useful in
PT diagnostics, forensics, gene mapping, identification of mutations
PT responsible for genetic disorders or other traits and to assess
PT biodiversity

PS Claim 20; SEQ ID NO 35173; 103bp; English.

XX The invention relates to isolated polynucleotide (I) and
XX polypeptide (II) sequences. (I) is useful as hybridisation probes,
XX polymerase chain reaction (PCR) primers, oligomers, and for chromosome
XX and gene mapping, and in recombinant production of (II). The
XX polynucleotides are also used in diagnostics as expressed sequence tags
XX for identifying expressed genes. (I) is useful in gene therapy techniques
XX to restore normal activity of (II) or to treat disease states involving
XX (II). (II) is useful for generating antibodies against it, detecting or
XX quantitating a polypeptide in tissue, as molecular weight markers and as
XX a food supplement. (II) and its binding partners are useful in medical
XX imaging of sites expressing (II). (I) and (II) are useful for treating
XX disorders involving aberrant protein expression or biological activity.
XX The polypeptide and polynucleotide sequences have applications in
XX diagnostics, forensics, gene mapping, identification of mutations in
XX responsible for genetic disorders or other traits to assess biodiversity
XX and to produce other types of data and products dependent on DNA and
XX amino acid sequences. ABG0010-ABG3037 represent novel human
XX diagnostic amino acid sequences of the invention.
XX Note: The sequence data for this patent did not appear in the printed
XX specification, but was obtained in electronic format directly from WIPO
XX at ftp.wipo.int/pub/published_pct_sequences.

XX Sequence 729 AA:

alignment_scores: Quality: 108.50 Length: 450
 Ratio: 0.512 Gaps: 19
Percent Similarity: 47.111 Percent Identity: 21.778

alignment_block:

US-09-303-518D-127/rev x ABG04814 ..

Align seg 1/1 to: ABG04814 from: 1 to: 729

1293 CCGCGGACAGAGGAGCGACAAAGCGAGTCTTCGTCGCAATTCGCA 1234

51 ProGlyProLysHisGlnProThrValArgIleuAlaIarGArgCysAr 67

1233 GCAACCGAATGCTGTC.....GGCTGTGGTATTCGCGGACGATTAAT 1190

67 GlnProArgAlaCysHisProAlaGlnAlaMetLeuLeuGlnGlnProL 84

1189 CGCGCAAAAGCAGGATAGCAGATGCTAGCGGATTTACGCGCTGTAA 1140
 ||| ||| ||| |||
84 euaArgProSerHisLeuGlnAlaArgHisLeuSerLeuPro..... 97
1139 GTACCAATCGGACCAATGCGCGGTCGCCACACCTTGACGCTGTGTA 1090
 ||| ||| ||| |||
98 ...ProLeuGlyLeuSerGlnSerGlyProProGlyLeuAlaProGlnPr 113
1089 CTGAGAGTTGTTGTTTTCAGAAATGCGGAGGTGTGTCGCGTATG 1040
 ||| ||| ||| |||
113 OLeuLeu**Leu.....GlnThrPro 121
1039 AGTATTGTCCGCTGC..... 1023
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121 SPHisPhalaGlyAlaIasProCysSerIleHisGlyThrSerHis** 137
1022GGCGCAACCCAGCGCAC.....AGCTTTTGTGCGCGCTTC 985
138 HisGlnGlyThrSerAsnProSerGlyArgAspGlyTyrGlnThrProSe 154
984 TTCGATACGGAATCTGATGTGTGTCGTCGCCAAATATACGCGCGC 935
 ||| ||| ||| |||
154 HisIleCysPro.....SerProAlaP 162
934 CT.....TGCTAATCGCGCGCTTCATACGAGACGGAATACAG 894
 ||| ||| ||| |||
162 rGlyGlnSerPheLeuPheGlyThrGlnAsnThrSerProSerSerPro 178
893 CGGTGTCTGCTCAACCAATTCGCGCGCAGTAATTCGATCTTCG 844
 ||| ||| ||| |||
179 AlaAlaProAlaAlaSerSerAlaProPheMetPheLysProIlePheTh 195
843 ACCCAAAACGTAACGACAGAGCGGTGTGTCGTCGAGAACGCCA 794
 ||| ||| ||| |||
195 rAlaProProLysSerGlyLysGlyLysProThr.....ProProGlyP 210
793 AAGCAATCAGCGCGTCGCTGTTCAGACGCGCTGTTCGAACAAAGCTCG 744
 ||| ||| ||| |||
210 rSerValThrAlaThrAla.....ProSerSerSerSerLeuPro 223
743 AAGCAATACATCTGATTAATGATGTCGAACAGGTTTGTTCGACC 694
 ||| ||| ||| |||
224ThrThrThrSerThrThrAlaPr 231
693 GACCGCTCATATGAATGATGTGCGTGCACCTCAACCG.....GCGC 650
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231 oThr.....PheGlnProValPheSers 239
649 GATCGCGCGCGCGAATTCATGTTGATGTTGGACGATTTTCAGAC 600
 ||| ||| ||| |||
239 erMetGlyProProAlaSer...ValProLeuProAlaProPhePheLys 254
599 GGCAGCTGTGCGCGAGCTGCTTACACACATGATTTTCGTCGTCGA 550
 ||| ||| ||| |||
255 GlnThrThrThrProAlaThrAlaProThr..... 264
549 ACGGTCATATACGACGACANACNTGTCGAATCMTGNCGCGCTTTCGA 500
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265ThrThrAlaProLeuP 270
499 TCACAACACAGAGGTGCGCGCAGGAGTGTGTCATTCGATTCGAG 450
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270 herThrGlyLeuAlaSerAlaThrSerAlaValAlaProIleThrSerAla 286
449 AAGATGGCGAAGCGCTCGGATCG.....ACGCGAGGATTTTGTCTGA 406
 ||| ||| ||| |||
287 SerProSerThrAspSerAlaSerLysProAlaPheGlyPheGlyLys 303
405 CGGACGNTACGAGGAGCAGTCCACAACCGGATTCGATTCGATTCG 356
 ||| ||| ||| |||
303 nSerValSerSerSerSerVal...SerThrThrThrSerThrAlaThrA 319
355 NNAATTCNTCGCGCTTAAGTTGCC.....AAGCTTCGCGCGCGGTG 312

```

319 1a1aSerGIInProPheLeuPheGlyAlaProGlnAlaSerAla1a1aSer 335
311 CGTGCAGACTCGATTTCGCGTGG.....CCTCAACGGC 277
336 PheIInProAlaMeCelSerIlePheGlnPheGlyLysProProAla1e 352
276 AATCAGACCGCAGTGAAGTACGCGCTTTTCGCGCATGATGAGCGGCGA 227
352 uProThrThrThrThrThrThrThrPheSerGlnSerLeuIsthrAla1v 369
226 TTTTGCCTGAACGCGCGCGTAACACACGACGCGCGATNCTTTTGTCT 177
369 alProThrAlaThrSerSerSerAlaAlaAspPheSerGlyPheGlySer 385
176 TCACACACACT...TGCGCTTTTTCAGCGATCGCGCTTCCTGACTT 130
386 ThrLeuAla1aThrSerAlaProAla1aThrSerSerGlnProThrLeuPh 402
129 CATCAGAGGCGCATACCGCATATTCCTCCCAAGCAGCAGCATTCG 80
402 eSerAsnThrSerThrProThrPheAsnIleProPheGlySerSerAla1 419
79 TAATACGGGCGCCCATAAATGACTTCCTCGGCTGCGCGCATGGGC 30
419 ySerProLeuProSerTyrProGlyAlaAsnProGlnProAlaPheGly 435

seq_name: /SIDSI/gcgdata/geneseq/geneseq-emb1/AA2001.DAT:ABB59201
seq_documentation_block:
ID ABB59201 standard; Protein; 746 AA.
XX
AC ABB59201;
XX
DX 26-MAR-2002 (first entry)
XX
DE Drosophila melanogaster polypeptide SEQ ID NO 4395.
XX
KW Drosophila; developmental biology; cell signalling; insecticide;
XX pharmaceutical.
XX
OS Drosophila melanogaster.
XX
PN MO200171042-A2.
XX
PD 27-SEP-2001.
XX
PF 23-MAR-2001; 2001WO-US09231.
XX
PR 23-MAR-2000; 2000US-191637P.
XX
PR 11-JUL-2000; 2000US-0614150.
XX
PA (PEKE ) PE CORP NY.
XX
PI Venter JC, Adams M, Li PMD, Myers EW;
XX
DR MPI: 2001-656860/75.
XX
DR N-PSDB; ABI03304.
XX
PT New isolated nucleic acid detection reagent for detecting 1000 or more
XX genes from Drosophila and for elucidating cell signalling and cell-cell
XX interactions -
XX
PS Disclosure: SEQ ID NO 4395; 21bp + Sequence Listing; English.
XX
CC The invention relates to an isolated nucleic acid detection reagent
XX capable of detecting 1000 or more genes from Drosophila. The invention is
XX useful in developmental biology and in elucidating cell signalling and
XX cell-cell interactions in higher eukaryotes for the development of
XX insecticides, therapeutics and pharmaceutical drugs. The invention
XX discloses genomic DNA sequences (AB16176-AB130511), expressed DNA
XX sequences (AB101840-AB16175) and the encoded proteins
XX (ABB57737-ABB72072).

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CC The sequence data for this patent did not form part of the printed
CC specification, but was obtained in electronic format directly from WIPO
CC at ftp.wipo.int/pub/Published_pcl_sequences.
XX
SQ Sequence 746 AA:

alignment_scores:
    quality: 107.00      length: 524
    ratio: 0.476         gaps: 21
    Percent similarity: 42.939      Percent identity: 20.229

alignment_block:
US-09-303-518D-127/rev x ABB59201 ..

Align seg 1/1 to: ABB59201 from: 1 to: 746

1334 TTCTCAANGGTTCCAGACCTTACGCAACAGGCGCGTATTCGATT 1285
||||| |||:||||| :|||:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|
53 PheSerHisGluSerThrThrLeuProThrSerAlaLeuPheLys...Le 68
1284 GCCCGGCGAGCAGACAGCTGCACAAACGAGGTTCTTCTGCAATTC 1236
| :|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|
68 uThrThrGluSerSerLeuSerSerSerSerSerSerSerSerThr 85
1235 .....AACCAACCAATCTGCGCG 1215
85 laHisIsthrThrAsnGluLeuLeuLysAsnValThrGlnSerTyr 101
1214 CTGTGCGATTCGCCAGCATTAATCGCGCAAAAGCAGGATGCGAGAT 1165
|||||:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|
102 LeuSerThrAlaLysProAlaProValArgThrThrSerAsnGlyThr 118
1164 GTCTAGCGGCATTACGCGCTCGTAAGTACCAATCGGCACCATGCGG 1115
|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|
118 uSerThr.....ThrArg 123
1114 CGCCACCGTTGAGGCGTGCCTGCACTGAAGAGTTGTTTCGAA 1065
|||||:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|
123 rgProProLeuAlaHisThrHisLysAsnSerHisTyr..... 136
1064 TGGCCGAGGCGTACGCGGTGATGAGATTTGTCGCGCGCGCAAC 1015
|||||:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|
137 .....ValValArgThrThrMetLysProSerAlaAlaProThrTh 150
1014 CCAGCGAAGACCTCTTGTGCGCGCTTTCGATACGCAATCTGAT 965
|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|
150 rThrIleThrProThrValLysProProArgArgArgThrThr... 165
964 TGTGTAGCGTCCCAATATATCGTCCGCGCTTGTGA..... 927
||:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|
166 .....AlaLysProValLeuThrThrAlaProIleValGluGlnArgIle 180
927 .....
181 GluThrThrSerValProThrLysPheValThrPheGlnIleValGluTh 197
926 ATCGCGCGCTTCATACCGAAGCG..... 903
197 rThrThrProAlaValThrGluProThrHisGlnGluThrThrTh 214
902 .....GAA 900
214 hrThrThrArgHisThrArgProThrGlySerSerSerValAlaSer 230
899 ATCAGCGGTTGTCTGCGTCAACCAATTCGCCAGATATTCGATAC 850
|||:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|
231 ArgThrThrSerLysThrThrThrThrThrThrThrThrThrTh 247
849 TTTTGCACCCAAACGATGACGAGCGCGTGTGACTGAGAAC 801
| ||||| |||:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|
247 rAlaAlaProProThrThrThrThrThrThrThrThrThrThrTh 264

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800 .....CCACCCAAAGCAATCACGGCTCG.....777
264 rgrProthrProleuargThrThrSerLysProHisLysThiHis 280
776 GTGTTACAGACGCGCTGTTCAAACAAAGTCGATGCGAATACATCTTG 727
281 SerSeratgargProAlaProAlaLysLysProValSerThrThr....295
726 ATAAATGATGGTCCAAACGGTTTGTGTTCACGACCGCGCTCAATGAAT 677
296 .....ThrThrThrAlaSerGlyThrGlnSerThrArg. 306
676 GAATGTGGTGGCACTCAACCGCGGATGCGGCGCCGATTCATGT 627
307 .....LysProAla.....AspSer 311
626 GTTTCGATGTTGGCAGCATTTTTCAGACGCGACGCTCGG.....CC 586
312 LeuAlaSerGlySerThrThrAlaSerGlyThrThrAlaThrThrLeuPr 328
585 AGCTGCTTACACACATGATTTTACGTCGCTCAACCGCTCAATACCA 536
328 AlaArg.....ArgProValThrGlnThrSerSerS 339
535 GCANNCNTGCTGTAATCNCGCGGCTGCTTGATCAAC.....492
339 erSerThrThrThrSerThrThrSerThrIleThrThrThyAlaPhe 355
492 .....492
356 LysAspGluLeuProLysAsnArgThrThrThrArgLeuProGluArgTh 372
491 .....ACAGG.....TCTG 482
372 rThrGlnAlaProArgProArgProLysProThrGlyLeuIleValLysV 389
481 CGCNAAGCGGATGGTGCATCGCATTCAGACAGATGCGAAGCGCTCG 432
389 alProAlaLeuLeuAlaGluProThrThrThrThrValThrLysGlySer 405
431 GCATCGACGCGCAGGAGATTGTGTAACGCGAGTACGACGCGCATCCA 382
406 SerSerSerSer.....SerSerSerPr 413
381 CAACCGCGATTGGATCAGATTGCNNNNNTCNCGCGCT.....339
413 oLysProPro.....AlaSerSerThrThrThrProLeuValThrL 426
338 ..AAGTTGCAACGCTTCGGGCGCTAGCGTTGCAACTCGATTGCTGC 291
426 yslLysLysProThrThrThrAsnAlaProThrThrThrThrThrThr 442
290 TTGCTTCAACGCAATCAACGACGCACTGAGTACGCGCTTTCGCGCG 241
443 ThrThrThrThrThrThrThrProLysProThrArgThrLysPr 459
240 ATGGAAGCGCGCATTTTGCCTCAACNGCGGCGTAACACACGCGCG 191
459 o.....ProThrThrThrThrThrThrThrLysA 470
190 GATNCCTTTTGTTCAAACAGCACTTGCGCTTTTTCAGCGGATGCGCT 141
470 lathThrThrThrThrLysAlaThrThrThrThrThrThrAlaLysPro 486
140 TCCTTGACTTTCATCNAGGCGCATACGCGCATATCTTCGCAAGCA 91
487 LeuIleThr.....ThrGluProProThrSerAlaProLeuTh 499
90 CGGACTTCGCTAATGACGGC 69
499 rThrThrThrLysLysThrGly 506
seq_name: /SIDSI/gcgsdata/geneseq/geneseq-emb1/AA2001.DAT:ABB64198

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seq_documentation_block:
ID   ABB64198 standard; Protein: 2406 AA.
XX
AC   ABB64198;
XX
DT   26-MAR-2002 (first entry)
XX
DE   Drosophila melanogaster polypeptide SEQ ID NO 19386.
KW   Drosophila; developmental biology; cell signalling; insecticide;
KW   pharmaceutical.
OS   Drosophila melanogaster.
XX
PN   WO200171042-A2.
XX
PD   27-SEP-2001.
XX
PF   23-MAR-2001; 2001WO-US09231.
XX
PR   23-MAR-2000; 2000US-191637P.
XX
PR   11-JUL-2000; 2000US-0614150.
XX
PA   (PEKE ) PE CORP NY.
PI   Venter JC, Adams M, Li PWD, Myers EW;
PI   WPI, 2001-656860/75.
DR   N-PSDB; ABL08301.
XX
PT   New isolated nucleic acid detection reagent for detecting 1000 or more
PT   genes from Drosophila and for elucidating cell signalling and cell-cell
PT   interactions -
XX
PS   Disclosure: SEQ ID NO 19386; 21pp + Sequence listing; English.
XX
CC   The invention relates to an isolated nucleic acid detection reagent
CC   capable of detecting 1000 or more genes from Drosophila. The invention is
CC   useful in developmental biology and in elucidating cell signalling and
CC   cell-cell interactions in higher eukaryotes for the development of
CC   insecticides, therapeutics and pharmaceutical drugs. The invention
CC   discloses genomic DNA sequences (ABLI01840-ABLI16175) and the encoded proteins
CC   sequences (ABBS7737-ABBT2072).
CC   (ABBS7737-ABBT2072).
CC   The sequence data for this patent did not form part of the printed
CC   specification, but was obtained in electronic format directly from WIPO
CC   at ftp.wipo.int/pub/published_pcl_sequences.
XX
SQ   Sequence 2406 AA;

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alignment_scores:

Quality	107.00	Length:	505
Ratio:	0.502	Gaps:	25
Percent Similarity:	42.178	Percent Identity:	20.594

alignment_block:

US-09-303-518D-127/rev x ABB64198 ..

Align seg 1/1 to: ABB64198 from: 1 to: 2406

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1298 CCNAT.....TCGATTTGGCGGCGCAGACGAGCT 1267
1266 rProLysGlyLeuThrHisGlySerGlyLeuProValLeuProVa 1282
1266 G.....CACAAGCGAGCT 1253
1283 lAlaThrProAsnLeuSerAsnLeuProThrGlnHisArgSer...S 1298
1252 CTTTCGTCGCAATTCGAAGCA.....CCC 1227
1298 erAspSerArgAsnSerArgLysSerProAlaSerLeuLysSerThrPro 1314

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1226 AATGCTGGCGCTGCGTATCG.....CGACGATTAAATCG..... 1188
1315 SerAsnIleGlyLeuAsnValSerMetAlaProThrIleArgSerIleTh 1331
1187 ..CGAAAAGCAGGAGTAGGAGATGTCTAGGCGCATTCAGCGCTGTAG 1139
1331 rProleuAsnAsnSerSerAlaIleSerSerGlyAlaSerGlnProVal 1348
1138 TACCAATCGGACCATAGCGCGGTGCGCACCGTTGACGCGTGTGTGTAAC 1089
1348 AlSerValValProSerAlaAsnSerThrAlaLeuSer..... 1360
1088 TTGAAGATTGTTGTTTTCAGGAATGCGCGGCTGTA.....CG 1048
1361 .....MetSerAsnProHisIleSerHisSerHisH 1371
1047 CGTATGAGATATTGTCGCGC.....TGCGGCG 1019
1371 sValProAlaThrAlaSerGlyAlaPheSerSerAlaAlaIleGlyT 1388
1018 CAACCCAGCCGACACAGCTCTTTG.....CTGCGGCTTCTTCGATACG 975
1388 hrSerThrProAsnSerGlyLeuSerThrLeuAlaValThrSerLeuSer 1404
974 GAAATGTGATGTGTAGCGCTCCCAATATTCGCGCGCTGTGTAAAT 925
1405 Thr.....SerAlaAlaPro..... 1409
924 CGCGCCCTCAATACCGACCGAATACAGCGGTG..... 888
1410 GlnProHisSerHisPheProGlnSerThrGlnMetLeuProGlnSer 1426
887 .....TTCGGCTCA 879
1426 LysnPheSerSerValSerHisLeuThrThrHisPheMetSerSer 1442
878 ACCAATTCGCCCGCAGTAAATTGCGACTTTCGACCCCAAAAGCGTAGC 829
1443 GlnAsnGlnProMetValArgCysGlySer.....ThrLeuTy 1455
828 CAAGAGCGCGTGTGTTGTTGACTTGAAGAACCCCAAGCATCAGCGCT 779
1455 rSerGlnSerSerAlaAlaThrAlaProProSerAlaAlaAla.... 1470
778 CGGTGTTCACAGCGCTGTGCAACAACAACATCCGATGCAATTCATCT 729
1471 .....AlaValSerAsnPheThrProSerVal..... 1479
728 TGATAATTGATGGTCCAAACGGTTTGTTCACACGACCGCGCATGAA 679
1480 .....LeuAlaValGlnSerLeuThrThrAlaValThrSerSer.... 1492
678 ATGAATGTGCGTCCACTCAAAACGCGCGGATGCGGCGCGCAATTCAT 629
1492 ..... 1492
628 GTGTTTCGATGTTCGACATTTTCAGACGACGAGTTCGCGCAGCTGCC 579
1493 .....SerSerSerProSerThr 1498
578 TTACACACATGGATTTCAGCTCGTC.....AAACG 547
1499 LeuSerSerSerValIleGlnIleValIleSerProIleGlnIleSerPr 1515
546 GCTCAATACGACANACNTCGTGAATTCNTGNCGCGCTCTTGTGATCA 497
1515 OCysAsnIleAspArgAspSerSerThrSerSerProAlaAsnAlaVal 1532
496 CAACCAACA.....GGGTCTGCC... 480
1532 AlThrThrCysAlaProThrThrProIleValSerSerGlySerAlaArg 1548

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479 .....CGNACCGGATTGGTGTG 463
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462 CATCGCATTTGACGAGATGCGGACGAGCGCTCGGCA..... 429
1565 nAlaAlaSerThrAlaArgSerSerCysAsnAlaIleSerProLeuSerI 1582
428 .....TGACGCGCAGGATTTGCTGACGACGAGCGGTTACGACGCGAGTC 384
1582 leProAlaThrAlaGlyIleHisValSerAlaThr..... 1593
383 CACAACCGGATGGATCAGATTGCNNCANNATTCCTGCGCCCTTAAGTT 334
1594 ...AsnProSerPheGlnSerSerSerThrPheProThrProLeuAlaPr 1609
333 TGCCAAACGCTTGCGGCGCGTAGCGCTTCGAACTGCAATTCGTCGTGCTT 284
1609 oProProSerSerProSerProAlaThrSerSerAlaAlaIleSerS 1626
283 CAACGCGCATCAACGACGCGTAGTACGCGCTTTTCGCGC..... 243
1626 erSerAla.....SerGlnPheAsnProAlaValSer 1636
242 CGATGATGCGCGCATTTTGCTGAAACNGCGCGGTAACACACGCGCC 193
1637 HisSerMetSerSerIleValThrThrAlaGlyAlaThrThrThrAl 1653
192 CGGATTCCTTTTGTCTTCAACAGCAGCTTGCGCTTTTTCGACGCGATCGC 143
1653 aSerSer.....ValThrGlnP 1659
142 CTTCCTTGACTTTTCATCNAGGGCGGCATACCGGCATATTCCTCCCAACG 93
1659 roSerValAlaAlaIleSerAsnProValThrAsnThrProHisProPhe 1675
92 AACGCGACTTCGCTA 78
1676 SerAlaGlnSerLeu 1680

seq_name: /SIDSL/gcgdata/geneseq/geneseqp-emb1/AA2001.DAT:ABG23390
seq_documentation_block:
ID ABG23390 standard; Protein; 1209 AA.
XX
AC ABG23390;
XX
DT 18-FEB-2002 (first entry)
XX
DE Novel human diagnostic protein #23381.
XX
KW Human; chromosome mapping; gene mapping; gene therapy; forensic;
KW food supplement; medical imaging; diagnostic; genetic disorder.
XX
OS Homo sapiens.
XX
PN WO2001/5067-A2.
XX
PD 11-OCT-2001.
XX
PF 30-MAR-2001; 2001WO-US08631.
XX
PR 31-MAR-2000; 2000US-0540217.
XX
PR 23-AUG-2000; 2000US-0649167.
XX
PA (HYSE-) HYSEQ INC.
XX
PI Drmanac RT, Liu C, Tang YT;
XX
WP1, 2001-639362/73.
XX
DR N-PSDB; AAS87577.
XX
PT New isolated polynucleotide and encoded polypeptides, useful in

```



```

912 6TALALysSerProLeuProSerTyrProGlyAlaAsnProGlnProAla 928
35 ATGGGC 30
929 pheGly 930

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seq_name: /SIDSL/gcgdata/geneseq/geneseqp-emb1/AA2001.DAT: AAG90820

seq_documentation_block:

ID AAG90820 standard; Protein; 1209 AA.

AC AAG90820;

DT 26-SEP-2001 (first entry)

DE C glutamicum protein fragment SEQ ID NO: 4574.

KW Coryneform bacterium; amino acid synthesis; vitamin; saccharide;

KW organic acid synthesis.

OS Corynebacterium glutamicum.

PN EPI108790-A2.

PD 20-JUN-2001.

PF 18-DEC-2000; 2000EP-0127688.

PR 16-DEC-1999; 99JP-0377484.

PR 07-APR-2000; 2000JP-0159162.

PR 03-AUG-2000; 2000JP-0280988.

PA (KXOW) KYOMA HAKKO KOGYO KK.

PI Nakagawa S, Mizoguchi H, Ando S, Hayashi M, Ochiai K, Yokoi H;

PI Tateishi N, Senoh A, Ikeda M, Ozaki A;

DR N-PSDB: AAH66039.

DR WPI: 2001-376931/40.

PS Claim 17, SEQ ID NO: 4574; 246pp + Sequence Listing: English.

The present invention provides a number of nucleotide and protein sequences from the Coryneform bacterium Corynebacterium glutamicum. These are useful for identifying the mutation point of a gene derived from a mutant of coryneform bacterium, measuring expression amount and analyzing the expression profile or expression pattern of a gene derived from coryneform bacterium, and identifying a homologue of a gene derived from coryneform bacterium. Coryneform bacteria are useful for producing amino acids, nucleic acids, vitamins, saccharides and organic acids, particularly L-lysine. The present sequence is a protein described in the exemplification of the invention.

Note: The sequence data for this patent did not form part of the printed specification, but was obtained in electronic format directly from the European Patent Office.

Sequence 1209 AA:

alignment_scores:

Quality: 106.50

Ratio: 0.431

Percent Similarity: 44.909

alignment_block:

US-09-303-518D-127 x AAG90820 ..

Align seg 1/1 to: AAG90820 from: 1 to: 1209

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49 GAGCAGTCATTTATGACGGCGCCGCTCATTCAGGATCGCGTTCCTGG 98
649 LysSerValProMetAspArgValIleIleGlyAspValGly...TyrGI 664
99 CGAAGAAATATGCCGATGCGCCCTNGATGAAA...GTCAAGGAAGCG 145
664 YLysThrGluValAlaValArgAlaIlePheLysAlaValGlnAspGly 681
146 ATGCCGTC.....AAAAAGCCCA 165
681 ysglnValAlaValAlaValProThrThrLeuAlaGlnGlnIle 697
166 GTGCTGTTGAGACAAAAGNAT.....CCGGCGTGATG... 201
698 SerThrPheGluGluArgMetThrGlyPheProValThrIleLysGly 714
202 .....TTTACCGCGCCN.....GTTTCAGGCMAA 226
714 userArgPheThrSerProAlaGluSerArgGluIleLeuSerGlyLeu 731
227 TCAGCGCC.....ATGCATCGCGCGAAGAGCGCTACTTCAGTCG 267
731 laAlaGlySerValAspIleValIleGlyThrHisArgLeuGlnThr 747
268 GTCGTC..... 273
748 GlyValGlnThrPryAsnLeuGlyLeuValIleValAspGlnGlnArg 764
274 .ATTGCCGTTGAAGCAGCAGCAAAATCGATTGCAAGCTACGCCCGCG 322
764 gPheGlyValGlnIleLysGlnIleLysAlaLeuArgThrHisVal 781
323 AAGCGTTGCAAACTTAAGCGCGGANGAANTNNGNCAATGTGATCCAA 372
781 spAlleu...ThrMetSerAlaThrProIleProArgThrLeuGlnMet 796
373 TCAGGTTTGTGACGTGCGTGCATCCGCTGACGAAATCCCTGC 422
797 Ser.....MetAlaGlyIleArgGlnMetThrThrMetLeuThrPro 811
423 CGTCATGCCGACGCGTTCGCCATCTTCGTCATATGCGATGACACCAATC 472
811 oGluAspArgHisProIleLeuThrGlyValGlyProGlyAspLysG 828
473 CGCTNCGGCA..... 483
828 InValAlaIleSerIleArgArgGlnLeuLeuArgAspGlnValPhe 844
484 .....GACCCGTGCTTGATCAAGAAGCCGANGATTTTCAG 524
845 PheIleHisAsnLysValAlaAspIleGlyLysAlaArgGlnIleArg 861
525 A.....CGANGTNTGCTGTATGACCGCTTTCAGCG 556
861 gAspLeuValProGluAlaArgValValAlaHisGlyGlnMetSerG 878
557 AGCGTAAATTCAT..... 570
878 lncIleuLeuGlnGlnThrValGlnGlyPheTrpAspArgGlyTyrAsp 894
571 .....GTGCTGAAGCA.....GTCGCGCAGACGTCGCTGCA 605
895 ValLeuValCysThrThrIleValGlnThrGlyLeuAspIleSerAsn 911
606 AATGCTGCCAATCGAAGAACATGAATTCGCGCGCGATCCGCGCG 655
911 asnThrIleuIleValGlu.....AsnIleHisHisMetG 923
656 GTTTAGTGGCAGCAC..... 672
923 lyeuSerGlnLeuHisGlnLeuArgGlyArgValGlyArgSerArgGln 939

```

```

673 .....ATTCAATTCATTGACCCGCGTGGTCAACAAACCCGTTTG 713
      ::::: 111 1111111111
940 ArgGlyTyrAlaTyrPheLeuTyrProLysGlyAlaThrLeuThrGluMe 956
      ::::: 111 1111111111
714 GACCATCAATATACAGATGTAATTCGATCCGACGTTTGTTCACACAG 763
      ::::: 111 1111111111
956 tSerTyrAspArgLeuAlaThrIleAlaGlnAsnAsnAspLeuGlyAlaG 973
      ::::: 111 1111111111
764 GCCGCTGCAACCCGAGCGCGTATTCCTTGGGTGGTTCACATCAAC 813
      ::::: 111 1111111111
973 LymetAlaValAlaMetLysAspLeuGlnMetArgGlyAla..... 986
      ::::: 111 1111111111
814 AAACACGCGCTCTGCGTACCGTTTGGGTGGTCCGAAAGTACGCAATAC 863
      ::::: 111 1111111111
987 .....GlyAsnValIleuGlyAlaGlnSerGlyHisIle 998
      ::::: 111 1111111111
864 TGGC.....GGCAATTCGTTG 880
      ::::: 111 1111111111
998 eAlaGlyValGlyPheAspLeuTyrValArgLeuValGlyGlnAlaValG 1015
      ::::: 111 1111111111
881 ACAGACACACCGCGTATTCGCGTTCGATTCGATTCGACGCGCGATT... 927
      ::::: 111 1111111111
1015 LualatYArgAlaLeuAlaAspGlyLysValAlaAspGlyThrValLys 1031
      ::::: 111 1111111111
928 .....ACACAGCGCGGCAC..... 942
      ::::: 111 1111111111
1032 GlyProLysGlnIleArgValAspLeuProValAspAlaHisIleProG 1048
      ::::: 111 1111111111
943 .GATATTTGGGACGCGTACACCAATCAGATTTCGTTATCCAGAAAGGCC 991
      ::::: 111 1111111111
1048 ulysTyrIle .AsnAlaGlnArgLeuArgLeuGlnIleTyrArgLysLeu 1064
      ::::: 111 1111111111
992 GCAGCAAGAGCTGTGGCTGGTGGTGGCGGACCGGACAAATACTCC 1041
      ::::: 111 1111111111
1065 AlaGlnSerGlnSerGlnValAspLeuArg .LeuAlaValGlnGlnMetG 1081
      ::::: 111 1111111111
1042 ATCAGCGGTACAGACCGTCCGCGATTCCTGAAACAAACCTCAACT 1091
      ::::: 111 1111111111
1081 LAspArgTyrGlyProIleProGlnGlnAlaGlnArgLeuAlaVal 1097
      ::::: 111 1111111111
1092 CACGACAGCGCTCAACGGTGGCGACCGCGCAT.....GGTGC 1129
      ::::: 111 1111111111
1098 SerArgLeuArgHisLeuMetArgGlnAlaHisLeuThrAspIleAlaVa 1114
      ::::: 111 1111111111
1130 CCATTGGTACTACGACGCGGTATCCGCTAGACATCTGCTACCC... 1176
      ::::: 111 1111111111
1114 Gln .GlyThrArgIleLysValHisProValAspLeuAlaAspSerGln 1130
      ::::: 111 1111111111
1177 ...CTGGCTTTCGCGGATTTATCGCGGCGATACCGACGCGCGCAAGC 1223
      ::::: 111 1111111111
1131 GlnValArgLeuArgLeuArgPheProGlnAlaThrTyrArgAla...Al 1146
      ::::: 111 1111111111
1224 ATTGGGTTCTTGGAATTGGACGAAGAACCTCGCTTGTGCACCTTCG 1273
      ::::: 111 1111111111
1146 aAlaLysAlaIleGlnLeu.....SerPheP 1155
      ::::: 111 1111111111
1274 TCTGCCCGGCGCAATACGAATANGCGCGCTTTCGCTAAGTG 1317
      ::::: 111 1111111111
1155 rLysThrGlyAsnLysValThrAspProLeuLeuArgAspVal 1169
      ::::: 111 1111111111

```

seq_name: /SID1/gcgdata/geneseq/geneseq-emb1/AA1999.DAT:AA04955

seq_documentation_block:

ID: AA04955 standard; Protein: 573 AA.

AC: AA04955;

DT: 06-JUL-1999 (first entry)

DE: Mycobacterium species protein sequence 41173.

KW: Secreted protein; Mycobacterium; primer; PCR; amplification; probe;

```

KW hybridisation; detection; vaccine; immunisation; infection.
XX Mycobacterium sp.
OS WO9909186-A2.
XX 25-FEB-1999.
XX 14-AUG-1998; 98WO-FR01813.
XX 11-SEP-1997; 97FR-0011325.
XX 14-AUG-1997; 97FR-0010404.
XX (INSP ) INST PASTEUR.
XX Gicquel B, Lim EM, Pelicic V, Portnoi D, Goguet de la Salmoniere Y;
XX Guineau A;
XX WPI: 1999-181045/15.
XX N-PSDB: AAX34206.
XX Mycobacterial DNA vectors containing reporter constructs - for
XX PT Identifying coding or promoter sequences involved in
XX PT infection-associated protein expression
XX PS Claim 32; Fig 41T; 309pp; French.
XX Sequences AA04742-Y05000 and AA07201-Y07204 represent secreted
XX CC proteins from various Mycobacterium species microorganisms. The
XX CC encoding nucleotide sequences can be used as primers and probes for
XX CC methods for detecting and identifying mycobacteria, especially belonging
XX CC to the M. tuberculosis complex. The encoded proteins can be used in
XX CC vaccines for immunisation against a bacterial or viral infection.
XX Sequence 573 AA:
SQ

```

alignment_scores:

Quality:	105.50	Length:	521
Ratio:	0.493	Gaps:	29
Percent Similarity:	41.075	Percent Identity:	24.568

alignment_block:

US-09-303-518d-127 x AA04955 ..

Align seg 1/1 to: AA04955 from: 1 to: 573

```

35 TCGCGGCGACGCGACCAAGTCAATTATACGCGCGCGATTA..... 79
    |||  ::  |||||  ::  |||  ::  |||||
45 SerThrSerThrArgSerThrGlySerMetCysSerArgSerLeuThrPr 61
    |||  |||  |||  |||  |||  |||  |||  |||  |||  |||
80 ...CCGAGTCGCGCTTGGCGGAGAAATATCCGCTATGCGCGCTN 125
    |||  |||  |||  |||  |||  |||  |||  |||  |||  |||
61 cIleProAlaSerThrCysSerPro.....CysValPro. 73
    ::::: 111 111 111 111 111 111 111 111 111 111
126 GATGAAGTCAGAGAGCGATGCCGTCAAAAAGGCGAAGTGTGTTG 175
    ::::: 111 111 111 111 111 111 111 111 111 111
74 ....ArgSer**LysLeuIleCysThrArgIleArgArgLeuThrPro 88
    ::::: 111 111 111 111 111 111 111 111 111 111
176 AAGACA..AAAAGNATCCGGCGCTGTACCG.....CG 210
    |||||  ::  |||||  |||  |||  |||  |||  |||  |||  |||
89 LysThrTrpArgGlyLeuArgProLeuSerArgProAlaArgValGlnAr 105
    ::::: 111 111 111 111 111 111 111 111 111 111
211 CCNGTTCAGGCAAAATCGCGCGCATTCATCCGCGGCGGCAAAAGCGCTACT 260
    ::::: 111 111 111 111 111 111 111 111 111 111
105 g.SerSerArgGlnLeuArgArgAspProValProAspProAlaArgAsp 121
    ::::: 111 111 111 111 111 111 111 111 111 111
261 TCAGTCGT...CGTATTCG.....CG 280
    |||  |||  |||||  |||  |||  |||  |||  |||  |||  |||
122 ArgArgGlyAspArgAspCysGlyArgArgLeuTrpArgGlyIleAlaG 138
    ::::: 111 111 111 111 111 111 111 111 111 111
281 TTGAAGGACAGCAGAAATCGAGTTTCAGACGCTACGCGCGCGAAGCGTTG 330
    |||  ::::: 111 111 111 111 111 111 111 111 111 111

```

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138 yelYargArgGlnArgProArgGlyArgValAlaAlaArgArgValG 155
331 GCAAACTTAAGCGGGANGAANTNNGCAATCTGATCCAAATCCGGTTT 380
155 LY.....ArgArgValAspGlyTTPArgGlyGlyAlaGlyAsnTyr 168
381 GTGAGCTGGCTCGGTANCCGTCCTCAGCAAAATCCCTGGCGG..... 425
169 ArgLeuGlyAlaAlaGlyArgArgArgSerArgArgProValAlaArgAlaAr 185
426 .....CGATCCGAGCGCGTTCGCCATCTTCGTCAATGCGATGAGAC 465
185 gclYValGlyArgCysGlyHisArgArgArg***Arg.....GlyGlyH 200
466 ACCAA.....TCGGCTNCGCGAGACGCTGTGCT 494
200 ISArgAlaGlyLysAspProArgThrAla***ArgAlaArgArgCysGly 216
495 TGTGATCAAGAGCGCGANGANTTCAGACGANGTNGCTGTGATTGA 544
217 ArgGlyArg.....ArgArgThrGlyProAlaGlySerAlaGly..ArgV 232
545 GCCGTTTGACCGAGCGTAATAATCCATGTGTGAAGCAGCGTGGCGAGAC 594
232 ALAlaLeuHisLeuArgGlyArgGlyThrCysProGlyGlyLeuArgThr 248
595 GTGGCGTGTGAATGCTGCCAATCGAAACATGATGATTCGGCGGCC 644
249 LeuAlaAlaArgValAla.....AspArgHisGlyTyrProThrPr 262
645 GCATCCGCGC.....GGTTGAGTGGCAGCGACACATTCAT 679
262 oArgProAlaAlaLeuArgGlyAspValSerValGlyMet***CysCys 279
680 TCATTGACCGGTC...GTGCAAAACAAAACCGTTTGACCATCAATTA. 725
279 erGlyGlyProValAlaGlySerThrGlnGlyLeuGly***ValGlyGly 295
726 ...TCAGATGTATTCGCATCGAGCTGTGTTGCACAGCGCGCTGCA 772
296 HisArgArgCysSerAla...ArgGlnLeuLeuArgThrArgPro..... 309
773 ACACCGAGCGCGTATTC.....TTGGTGGTGTGCA 807
310 .HisArgArgArgCysArgArgGlySerArgGlyLeuGlyAla 336
808 GTCAACAACACGCGCTTCGCGTACCGTTGGGTGCGAAATATCGCA 857
326 er***ProAspArgAspLeuGlyAlaArgPheArgAspGlnArgGlyLeuAla 342
858 AATTACTGC.....GGCGAATTGCTTGACCGCAGACACC 892
343 GlyArgTrpLeuAspAlaGlyProArgArgAlaGlyGlyArgArgArg 359
893 GCGGTATTCGG.....TTGC 909
359 gATGcysArgArgAlaValaArgArgGlyArgLeuArgAlaAlaThrG 376
910 GTATTGACGCGCGGATTACACAGG..... 935
376 LysArgArgArgAspTrpGlyArgArgTyrGlnCysProProAlaGly 392
935 ..... 935
393 AlaglyArgGlyArgHisArgArgArgAlaArgAspGlyAlaAlaGlnTr 409
936 .....CGGCACGATTTATTTGGAGCGTACCAACATGAGATTTC 975
409 pleuGlyArgArgArgThrGlyGlyArgValTyrArgGlyAsp...A 425
976 GTTATCGAAGAGCGCGCAGCAAGACTGTTGCGCTGGCTGGCCCGCA 1025
425 rgluGlyArgArgArgGlyThrArgAlaAspArgIleAspGlyAlaGly 441

```

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1026 GCGGACAAATA..... 1037
442 ValGlyArgAlaGlyArgAla***ArgGlyProProGlyArgArgArg 458
1038 .CTCCATCAGCGGTACGACCGCTGGCCATTTCTGTAATAAACAATCTTC 1086
458 gIeuGlnHisGlyProCysArgArgCysPheProAlaArgIleGlyAlaH 475
1087 AAGTTCAGCAGCAGC..... 1100
475 isCysHisProLysGlyAlaAspLeuGlyArgTyrLeuGlnAlaGlyArg 491
1101 CGTCAACGCTGGCGCGCGCCATGTCGCGATTTGCTACTTACGAGCGG 1150
492 ArgSerGlyTyrArgGlyArgArgGlyAlaAsp..... 502
1151 TAATGCCCTAGACATCTGCTACCCCTGTTTGGCGGATTAATCTGTC 1200
503 .....ArgArgA 505
1201 GCGGATACGAGCAG.....CGGCACGATTTGGTGTGGAAT 1241
505 rArgCysArgArgGlyGlyHisArgSerGlyArgProValAlaGlyIle 521
1242 GGACGAGA 1250
522 GlyArgArg 524

```

seq_name: /SIDS1/gcgdata/geneseq/geneseqp-emb1/AA1999.DAT:AAW89449

seq_documentation_block:
ID AAW89449 standard; Protein; 372 AA.

AC AAW89449;

DT 18-MAR-1999 (first entry)

DE A gida2 polypeptide fragment.

KW gida2; Staphylococcus aureus WCUH29; bacterial infection;

KW Helicobacter pylori infection; Cancer; ulcer; gastritis; antibacterial;

OS wound treatment; Bacterial adhesion; matrix protein.

Staphylococcus aureus.

EP889131-A2.

30-JUN-1998; 98EP-0305203.

01-JUL-1997; 97US-0051380.

(SMIK) SMITHKLINE BEECHAM CORP.

(SMIK) SMITHKLINE BEECHAM PLC.

Burnham M, Deboeck CM, Kallender H, Lenox AL, Mooney JL;

Palmer LM, Zhong Y;

WPI; 1999-062662/06.

N-PDB; AAW82085.

New isolated gida2 polypeptide from Staphylococcus aureus - used to

diagnose, treat and prevent bacterial infections e.g. S. aureus and

H. pylori, related cancers, ulcers and gastritis and to prevent

adhesion of bacteria to matrix proteins

Claim 1; Page 6; 41pp; English.

The present sequence represents a gida2 protein fragment of

Staphylococcus aureus WCUH29. The gida2 proteins and nucleic acids

are used to treat conditions requiring increased activity or expression

alignment_scores:
 Quality: 105.00 Length: 272
 Ratio: 0.784 Gaps: 13
 Percent Similarity: 49.265 Percent Identity: 21.324

alignment_block:

US-09-303-518d-127 x AAW89448 ..

Align seg 1/1 to: AAW89448 from: 1 to: 435

```

331 GGAACCTTAAGCGCGGANGAANTNNGNCAATCTGATCATCCGGTTT 380
|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||
10 AAGlyLeuAlaGlySerGluAlaAlaTyrGlnLeuAlaGluArgGlyI 26
381 GTGACTCGCGTG...CGTANCCGCTGCAGCAAAATCCCTCCGCG 427
: : : : : : : : : : : : : : : : : : : : : : : : :
26 elysValasnLeuIleGluMetAlaProValIysGlnhrProAlaHis 43
428 ATGCCGACCCGCTGCCATCTTCGTC.....AAWGGATGACACCAAT 471
|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||
43 IsthAspIysPheAlaGluLeuValCysSerAsnSerLeuArgGlyAsn 59
472 CCGCTNCGCGGAGACCCGTGCTGTATGATCAAGACCGCGANGATT 521
|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||
60 AlaLeu...ThrAsnGlyValGlyValLeuLysGlu.....GluMe 72
522 CAGACGANGTTCGCTGTATGACCGCTTGACCGACGCTAAATTCATG 571
|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||
72 TArgArgLeuAsnSerIleIleGluAlaAlaAspLys.....85
572 TGTGTAAAGCAGCTGCGCAGACGCTGCTGAAAATGTCGCCACATC 621
|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||
86 .....AlaArgValProAlaGlyAlaAlaLeuAlaVal 96
622 GAAACACATGAATTCGGCGCCGCGCATCCGCGGTTTGAAGTGGACGCA 671
: : : : : : : : : : : : : : : : : : : : : : : : :
97 AspArgHisAspSerGly.....103
672 CATTCATTTCATGACCGCGCTGCTGCACAAACACCGTTGGACATCA 721
|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||
104 ....TyrIleThrGlnThrLeuLysAsnHisGluAsnIleThrValIle 119
722 ATTATCAAGATGTAATGCGCATCGA.....CGTTGTTGCAACA 762
|| : : : : : : : : : : : : : : : : : : : : : : :
119 sn...GluGluIleAsnAlaIleProAspLysTyrThrIleIleAlaThr 134
763 GCGCGTCTGACACCGAG.....CGCGTATTCCTTGGGTG 800
||| : : : : : : : : : : : : : : : : : : : : : :
135 GlyProLeuThrThrGlnThrLeuAlaGlnGluIleValAspIleThrG 151
801 TTCTCAAGTCAACAAACACCGCTCTGCTACCGTTTGGGTGCGAAG 850
|| : : : : : : : : : : : : : : : : : : : : : :
151 LysAsp.....GlnLeuTyrPheTyrAspAla 161
851 TATCGCAAAATTACTCGCGCAATGTTGACGACAGACACCGCGGATT 900
|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||
161 IalAlaProIleIleGluLysGlnSerIleAspMetAspLysValIleu 177
901 .....TCGGTTCGATTTCAGACGCGCGGATTC 929
|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||
178 LysSerArgTyrAspLysGlyGluAlaIleTyrLeuAsnCysProMet 194
930 ACAAGGCGGCGACGATTATTGGAGCGTACCAATCGATT..... 972
|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||
194 rGlu.....AspGluPheAsnArgPheTyrAspAlaValLeuGlu 208
972 ..... 972
208 IagIuValAlaProValAsnSerPheGluLysGlnLysTyrPheGluGly 224
973 .....TCGCTTATCGAAGAGCGCGACGACCAAGAGCTGTT 1007
|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||
225 CysMetProPheGluValMetAlaGluArgGlyArgLysThrLeuLeuPh 241

```

1008 CGGCTGGGTGCGCCG 1023
 |||||:|||||
 241 eGlyProMetLysPro 246

seq_name: /STDS1/gcgdata/geneseq/geneseq-emb1/AA1997.DAT.AAW13502

seq_documentation_block:

ID AAW13502 standard; protein: 1645 AA.

AC AAW13502;

DE 30-JAN-1998 (first entry)

DE B. pertussis adenylcyclase-haemolysin mutant delta-cla.

KW Mutant; Bordetella pertussis; adenylcyclase; haemolysin; wild type;

KW deletion; induction; antibody; Bordetella parapertussis; vaccine; human;

OS Bordetella bronchiseptica; veterinary.

OS Bordetella pertussis.

OS Synthetic.

FT Key

FT Region

FT Modified site

FT 919..924

FT /note="Lys at position 922 is modified by addition of

FT a fatty acid"

FT FR2736064-A1.

FT 03-JAN-1997.

FT 30-JUN-1995; 95FR-0007945.

FT 30-JUN-1995; 95FR-0007945.

FT (INSP) INST PASTEUR.

FT Betson F, Guiso N, Sebo P;

FT WPI; 1997-111454/11.

FT New immunogenic fragments of Bordetella adenylcyclase haemolysin

FT and related nucleic acid and antibodies, for use in vaccines and

FT immunotherapy

PS Claim 6; Page -: 55pp; French.

CC This is the sequence of a novel mutant, designated delta-cla, derived

CC from the Bordetella pertussis adenylcyclase-haemolysin (AC-Hly) in which

CC amino acids 827-887 of the wild type sequence have been deleted by

CC genetic engineering. The novel protein is able to induce production

CC of protective antibodies against the Bordetella species pertussis,

CC parapertussis and/or bronchiseptica, especially in protective vaccines

CC for human or veterinary use.

CC Note: this sequence is not given in the specification but is generated

CC from the wild type sequence.

CC Sequence 1645 AA;

alignment_scores:

Quality: 104.50 Length: 455
 Ratio: 0.522 Gaps: 24
 Percent Similarity: 43.956 Percent Identity: 21.538

alignment_block:

US-09-303-518D-127 x AAW13502 ..

Align seg 1/1 to: AAW13502 from: 1 to: 1645

XX Disclosure; Fig. 5; 43pp; English.

PS The protein is encoded by the third reading frame of HIV-2 SBL/ISY, a
XX proviral clone of HIV-2.
CC (Note: Revised entry submitted to correct the patent number format of
CC US Government-owned NIS applications to prevent clashes with ongoing US
CC granted patent numbers. For further information please visit the Derwent
CC web site at www.derwent.com/dwpl/updates/nis_us.html.)
XX

Sequence 3080 AA:

alignment_scores:
Quality: 104.50 Length: 433
Ratio: 0.562 Gaps: 25
Percent Similarity: 42.956 Percent Identity: 21.940

alignment_block:

US-09-303-518D-127 x AAP93285 ..

Align seg 1/1 to: AAP93285 from: 1 to: 3080

```

195 CGTGTGTTTACGGCCGNGTTTCAGCAAAATCGCCGATCATCGCG 244
    ||| :|: ||| ||| :|: ||| :|: ||| :|: ||| :|:
256 ArgHisLeuValLeuAlaArgArgGluSerGluArgGlyArgSerly 272
    :|: :|: :|: :|: :|: :|: :|: :|: :|: :|: :|:
245 GCGAAAGCGCGT...ACTTACGCGT...CGTATGCGCGTTGAA 285
    :|: :|: :|: :|: :|: :|: :|: :|: :|: :|: :|:
272 slyAsnSerArgGluThrSerSerGlyArgAsnArgAsnGlyGlu 289
    :|: :|: :|: :|: :|: :|: :|: :|: :|: :|: :|:
286 GCGAAGCAAAATCGAGTTCGACCTCGCGCCGAGCGTTGGCAA 335
    :|: :|: :|: :|: :|: :|: :|: :|: :|: :|: :|:
289 snAlaLySTyLysThrAsnSerThrThrArgGluArg...Glylys 303
    :|: :|: :|: :|: :|: :|: :|: :|: :|: :|: :|:
336 CTTAAGCGCGGANGAANTNNGNCATCTGATCCAAATCGGTTGTGA 385
    ||| ||| :|: |||
304 LeuProArgAlaThrAsnArgArgGln..... 312
    :|: :|: :|: :|: :|: :|: :|: :|: :|: :|: :|:
386 CTGCGGCTGCTANCGCTCCGTTACGAAATCCCTCGCGTCGATCCGAG 435
    ||| ||| :|: ||| :|: ||| :|: ||| :|: ||| :|:
313 .....LeuCysProSerAlaAlaGluSerProAsnProLysCysLeug 327
    :|: :|: :|: :|: :|: :|: :|: :|: :|: :|: :|:
436 CC...GTTGCCATCTTCGTCATGAGATGACACCAATCCGCTGCC... 479
    :|: :|: :|: :|: :|: :|: :|: :|: :|: :|: :|:
327 LyysValSerArgGlyGlyValArgGlyArgSerSerAlaGlyLe 343
    :|: :|: :|: :|: :|: :|: :|: :|: :|: :|: :|:
480 ..... 481
    :|: :|: :|: :|: :|: :|: :|: :|: :|: :|: :|:
344 SerGlyThrLeuArgArgLeuHisAlaLeuTySerAsnAlaLeuGly 360
    :|: :|: :|: :|: :|: :|: :|: :|: :|: :|: :|:
482 CAGACCCGTGGTGTGATCAAGAAAGCGCGANGATTTCAAGACGANGT 531
    ||| ||| :|: ||| :|: ||| :|: ||| :|: ||| :|:
360 yArgProSerSerSerAspAlaAsnAsnGlnArgAsnTyArgSerSera 377
    :|: :|: :|: :|: :|: :|: :|: :|: :|: :|: :|:
532 NTGCTGATTTGAGCCGTTTGACGAGCGTAAATCCATGTCGTAAAGC 581
    |||
377 rglengly..... 379
    :|: :|: :|: :|: :|: :|: :|: :|: :|: :|: :|:
582 AGCTGGCGAGACGCTCGTGAATGTCGCAACATCGAACAACATG 631
    ||| |||
380 .....CysThrThrSerAsnThr.. 385
    :|: :|: :|: :|: :|: :|: :|: :|: :|: :|: :|:
632 AATTGGCGCGCGCATCCGCGGTTTGAGT.....GCGACGACATTT 675
    ||| ||| :|: ||| :|: ||| :|: ||| :|: ||| :|:
386 .....ArgProLeuThr..SerGlyAlaAlaGlnArgSerThrArgIle 399
    :|: :|: :|: :|: :|: :|: :|: :|: :|: :|: :|:
676 CATTTTCATTGAGCGGTCGTCGCAACAAAACGTTTGACCATCATTA 725
    |||
400 HisSerArgAspAsnLysHisSerArgGlyThr..... 410
    :|: :|: :|: :|: :|: :|: :|: :|: :|: :|: :|:
726 TCAGATGTAATTCATCGCATCGGCGCTTTGTTGCAACAGCGCGTCTGAACA 775
    ||| :|: :|: :|: :|: :|: :|: :|: :|: :|: :|: :|:

```

```

411 ....AspArgMetAspValAlaArgLysSerCysThrSerArgLysHisL 426
776 CCGAG.....CGCGGATGCGTTTG 795
    |||
426 euGluMetAspProAspArgThrAlaGluValCysGlnAsnValGlnSer 442
796 GGTGGTTCCTCAAGTCAACAAACACGCTTCCTGACGCTT..... 837
    :|: :|: :|: :|: :|: :|: :|: :|: :|: :|: :|:
443 AsnGlnHisSerArgHisLysThrArgThrLysArgValValProLysLe 459
838 .....TTGGTGCAGAAATATCGCAATTAATCTGCGG 868
459 uCysGlyTLeuGlnLysLeuLysGlyArgThrAspArgCysSerSerg 476
869 GCGAATGTTGTCAGCAGAC.....AACCGCGATTT 900
    ||| ||| |||
476 LngLLeuAspAspProAspAlaAlaSerAlaAlaGluProArgLeuVal 492
901 TCCGGTTCGCTATTGAACGCGCGATTCACAAAGCGCGACAGAT.... 945
    |||
493 Ser.....ThrLysGlyThrArgAspGluSe 501
946 ..TATTGGACGCTACACATACATTCGCTTATTCGAAGAGCGCGCA 994
    ||| ||| ||| :|: ||| :|: ||| :|: ||| :|:
501 rTyThrLeuArgArgAspAlaAsnArgLeuSer.....ArgA 513
995 GCAAAAGCTGTTCCGCTGGCTGGCTGGCGCGACGC.....GACAA 1034
    :|: :|: :|: :|: :|: :|: :|: :|: :|: :|: :|:
513 sParG.....TrpThrArgProGlnGlyGlnThrAsnGlyArg 525
1035 ATACTCATACGCGGTACGACGCGCTCGGCCATTTCT..... 1070
526 SerLeuLysArgGlyHisAlaThrSerProTyProLysCysSerSerg 542
1071 ....GAAAA...CAACTTTCAGATTACGACAGCGT..... 1103
    ||| ||| :|: ||| :|: ||| :|: ||| :|: ||| :|:
542 oThrGlnLysGlyAsnValLeuGlnLeuThrLysGlyArgAlaLeuGlyL 559
1104 .....CAACGCTGG.....CGA 1115
559 ySThrMetProSerAlaLysThrGlyLeuGluMetArgValArg 575
1116 CCGCGCGCATGCT.....GCGATGCTGACTTACGACG..... 1148
    :|: :|: :|: :|: :|: :|: :|: :|: :|: :|: :|:
576 ThrHisHisGlyLysLeuProArgThrGlyThrPhePheArgAlaTrpH 592
1149 .....CGTAATGCCGCTAGACATCTGCTACCGCTGTTTGGCGGAT 1191
    :|: :|: :|: :|: :|: :|: :|: :|: :|: :|: :|:
592 rMetGlyLysGlnAlaProGlnLeuProArgGlyProLysPheAlaGlyA 609
1192 TTAATCGTCGCGGATACGACGCGCGGACGATTTGGTGTGGA 1238
    ||| :|: :|: :|: :|: :|: :|: :|: :|: :|: :|:
609 LaAsnThrAsnSerThrProAsnGlySerSerSerGlyProThrGly 624
seq_name: /SIDS1/gcgdata/geneseq/emb1/AA2001.DAT:AAU34397
seq_documentation_block:
ID AAU34397 standard; Protein; 433 AA.
XX AAU34397;
XX
XX 14-FEB-2002 (first entry)
XX
XX Staphylococcus aureus cellular proliferation protein #673.
XX
XX Antisense; prokaryotic cellular proliferation protein;
XX antibiotic; antibacterial; drug design.
XX
XX Staphylococcus aureus.
XX
XX MO200170955-A2.
XX
XX 27-SEP-2001.

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```

XX 21-MAR-2001; 2001WO-US09180.
PF
XX
PR 21-MAR-2000; 2000US-191078P.
PR 23-MAY-2000; 2000US-206848P.
PR 26-MAY-2000; 2000US-207727P.
PR 23-OCT-2000; 2000US-242578P.
PR 27-NOV-2000; 2000US-253625P.
PR 22-DEC-2000; 2000US-257931P.
PR 16-FEB-2001; 2001US-269308P.
XX
XX (ELIT- ) ELITRA PHARM INC.
XX
XX Haselbeck R, Ohlsen KL, Zyskind JW, Wall D, Trawick JD, Carr GJ,
XX Yamamoto RT, Xu HH;
XX
XX WPI: 2001-611495/70.
XX N-PSDB: AASS2256.
XX
XX New polynucleotides for the identification and development of
XX antibiotics, comprise sequences of antisense nucleic acids -
XX
XX Example 3; Seq ID No 5893; 511pp; English.
XX
XX The invention relates to antisense inhibitors of genes essential to
XX prokaryotic cellular proliferation, their use in identifying the
XX genes, their use in the discovery of novel antibiotics, the essential
XX genes themselves and the encoded proteins. The prokaryotes used are
XX Escherichia coli, Staphylococcus aureus, Salmonella typhi, Klebsiella
XX pneumoniae, Pseudomonas aeruginosa and Enterococcus faecalis. The
XX invention is also useful for the identification of potential new targets
XX for antibiotic development. The antisense nucleic acids can also be used
XX to identify proteins used in proliferation, to express these proteins,
XX and to obtain antibodies capable of binding to the expressed proteins.
XX The proteins can be used to screen compounds in rational drug discovery
XX programmes. The antisense nucleic acid sequence is also useful to screen
XX for homologous nucleic acids which are required for cell proliferation in
XX a wide variety of organisms. The present sequence represents an
XX essential prokaryotic cellular proliferation protein.
XX
XX Note: The sequence data for this patent did not form part
XX of the printed specification, but was obtained in electronic
XX format directly from WIPO at
XX ftp.wipo.int/pub/published_pcl_sequences.
XX
XX Sequence 433 AA:
SQ

```

```

alignment_scores:
    Quality: 104.00      Length: 272
    Ratio: 0.776        Gaps: 13
    Percent Similarity: 49.265    Percent Identity: 21.324

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alignment_block:

US-09-303-518D-127 x AAU34397 ..

Align seg 1/1 to: AAU34397 from: 1 to: 433

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331 GCAACTTAAAGCGCGAAGTANNNGCAGTATGATCCAGTCCGGTT 380
    |||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
8  AlAGlYleuAlAGlYserGIuAlAlAlAYrGIInleuAlAGlAlrGIYl 24
381 GTGACATGCGCTG...CGTANCCGTCGCTTACAGCAAAATCCCTGCCGTCG 427
    :   :   :   :   :   :   :   :   :   :   :   :   :   :
24 elYsValAsnleuileGluMetArProValLYsGIInrProAlAnISH 41
428 ATGCCGACCGCTTGGCATTCTGTC.....AATGCATGACACACCAT 471
    :   :   :   :   :   :   :   :   :   :   :   :   :   :
41 IsrInAspLYsPheAlAGlYleuValCYsSerAsnSerleuAlrGlyAsn 57
472 CCGCTNGGCGACCGCTGTGTGTGATCAAGAAAGCCGCGCAGCATTT 521
    ||| :   :   :   ||| |||:|||||:|||||:|||||:|||||
58 AlAlleu...ThrAsnGlyValAGlYAlleuLYsGlu.....GluHe 70

```

```

522 CAGACGANGTTCGTGTATTTAGCCGTTTACCGAGCGTAAATTCATG 571
    |||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||
70 tArGArleuAsnserlellelleGluAlAlAlAspLYs..... 83
572 TGTGTAAAGCAGCTGCGCAGACGTCGCCGCTTCAAAATCTGCCAATC 621
    |||:|||||:|||||:|||||:|||||:|||||:|||||:|||||
84 .....AlArGValProAlAGlYAlleuAlAlVal 94
622 GAAACATGATTCGGCGGCCGCCGATCCGCCGTTTGTAGTGCACGCA 671
    :   :   :   :   :   :   :   :   :   :   :   :   :   :
95 AspArGHisAspPheSerGly..... 101
672 CATTCATTTCAATGACCGCTGCTGTCGCAACAAACCGTTTGACCATCA 721
    :   :   :   :   :   :   :   :   :   :   :   :   :   :
102 ....TyrlePheGluAsnleuLYsAsnHisGluAsnlelleValille 117
722 ATATCAAGATGTAAATTCGCATCGGA.....CGTTGTTTGCACGA 762
    || :   :   :   :   :   :   :   :   :   :   :   :   :   :
117 sn...GluGluileAsnAlalleProAspGlyTyrThrillelleAlaTr 132
763 GCGCGCTGACACCGCAG.....CGCGTATGCTTTGGGTGG 800
    |||:|||||:|||||:|||||:|||||:|||||:|||||:|||||
133 GylProleuTrnTrnGluThrleuAlAGlYAlleuValAspPheTrnG 149
801 TTCTCAAGTCAACAAACACCGCTTCCGTACCGCTTTGGTGCAGAG 850
    :   :   :   :   :   :   :   :   :   :   :   :   :   :
149 YLYsAsp.....GluLeuTyrPheTyrAspAlAl 159
851 TATGCCAAATTAATTCGGCGCATTTGGTACGACGACCAACCGGTGAT 900
    :   :   :   :   :   :   :   :   :   :   :   :   :   :
159 lAlAlProillelleGluLYsGluSerlleAspMetAspLYsValTyrLeu 175
901 .....TCCGTTTGGTATTTGAACGCGCGGATTC 929
    :   :   :   :   :   :   :   :   :   :   :   :   :   :
176 LYsSerArGTYrAspLYsGlyGluAlAlAlTyrleuAsnCYsPrometh 192
930 ACAGAGCGCGCAGCTTATTGGACGCTACCAACATCAAGATT..... 972
    :   :   :   :   :   :   :   :   :   :   :   :   :   :
192 rGlu.....AspGluPheAsnArGpHeTyrAspAlAlValleuGlu 206
972 ..... 972
206 lAGlYAlAlProValAsnSerPheGluLYsGluLYsTyrPheGluGly 222
973 .....TCCGTTATTCAGAGAGCGCGCAGCAAAAGACTGT 1007
    :   :   :   :   :   :   :   :   :   :   :   :   :   :
223 CYsMetProPheGluValMetAlAGlYArGlyTrnleuLeuPh 239
1008 CCGCTGCGTTCGCCGCG 1023
    ||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||
239 eGlyProMetLYsPro 244
seq_name: /SIDS1/gcgdata/geneseq/geneseq-emb1/AA1997.DAT:AAW18663
seq_documentation_block:
ID AAW18663 standard; Protein: 387 AA.
XX
XX AAW18663;
XX
XX 24-JUL-1997 (first entry)
XX
XX Fragmented human NF-H gene +2 frameshift mutant product.
XX
XX Frameshift mutation product: GAGA motif, somatic mutation; diagnosis;
XX detection; antibody; probe; cancer; neoplasia; neurodegenerative;
XX Parkinson's; Alzheimer's disease; Pick's; Huntington's disease;
XX Down's syndrome; frontal lobe dementia; progressive supranuclear palsy;
XX PSP; amyotrophic lateral sclerosis; multiple sclerosis; MS;
XX cardiovascular; rheumatoid arthritis.
XX
XX Homo sapiens.
XX
XX OS
XX FH
XX Key Location/Qualifiers

```



```
1009 GCGTGGTGGCGCGCGGACCAATACATCCATCGCGGTACGACCT 1058
|| .....|||
317 rgarYserAlaAlaThrCysGlyAla..... 325
1059 CGGCGATTTCCTGAAAAACAACCTTCAAGTTACAGCAGCCGTCACG 1108
|| .....|||
326 ..... 330
1109 GTGGCGACCGCGCATGTGCGCATTTAGTACGAGCGCGTAATCCG 1158
||| .....|||
330 gtrpAlaSer.....CysSerAlaArgSerArg 340
1159 CTACACATCCGCTACCTGCTTTTCGGGATTATGTCGCGCGTAC 1208
|| .....|||
340 larpAlaPro.....ArgArgArgArgCys 348
1209 CGACAGCGCGCGACGATGGTGGTGTGGAATTGACGAGAGA...CC 1255
||| .....|||
349 ArgProArgArgAlaThrPro***SerAlaThr***ArgArgArgCysAl 365
1256 TCCTTTGTGCAG 1268
||| .....|||
365 aArgPheAlaArg 369
seq_name: /SIDSL/gcgdata/geneseq/geneseq-emb1/AA2000.DAT:AA15935
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seq_documentation_block:
ID   AAB15935 standard; Protein; 740 AA.
XX
AC   AAB15935;
XX
DT   05-OCT-2000 (first entry)
XX
DE   E. coli proliferation associated protein sequence SEQ ID NO:292.
XX
KW   Escherichia coli; E. coli; proliferation; inhibition; screening;
KM   antimicrobial; bacterial growth; antisense therapy; antibacterial.
XX
OS   Escherichia coli.
XX
PN   WO200044906-A2.
XX
PD   03-AUG-2000.
XX
PF   27-JAN-2000; 2000MO-US02200.
XX
PR   27-JAN-1999; 9905-0117405.
XX
PA   (ELITR-) ELITRA PHARM INC.
XX
PI   Zyskind J, Ohlsen KL, Trawick J, Forsyth RA, Froelich JM, Carr GJ;
PI   Yamamoto KT, Xu HH;
XX
DR   WPI; 2000-514822/46.
XX
DR   N-PSDB; AAB65940.
XX
PT   Novel polynucleotides and polypeptides associated with microorganism
PT   proliferation, used to identify inhibitors of bacterial growth and
PT   proliferation, for use in antisense therapy -
XX
PS   Claim 11; Page 217-219; 316pp; English.
XX
CC   AAB65809 to AAB65889 and AAB66058 to AAB66138 represent nucleotide
CC   sequences derived from Escherichia coli which inhibit E. coli
CC   proliferation. AAB65890 to AAB66055 and AAB15886 to AAB16040 represent
CC   nucleotide and protein sequences associated with E. coli proliferation.
CC   AAB66056 and AAB66057 represent sequences used for sequencing E. coli
CC   proliferation inhibiting nucleotide inserts in an example from the
CC   present invention. Methods from the present invention can be used to
CC   identify a proliferation-required gene in a microorganism, by contacting
CC   a microorganism with a proliferation-required gene actively inhibitory
CC   nucleic acid identified in another organism, and determining if
```

CC inhibition occurs in the second microorganism. The nucleic acid sequences
CC identified as being required for bacterial growth and proliferation, can
CC be used for antisense therapy for killing bacteria.

XX Sequence 740 AA;

alignment_scores: Length: 324
 Quality: 103.00 Gaps: 13
 Ratio: 0.665 Percent Identity: 21.914
Percent Similarity: 47.840 Percent Identity: 21.914

alignment_block:

US-09-303-518D-127 x AAB15935 ..

Align seg 1/1 to: AAB15935 from: 1 to: 740

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100 GAAGAAATATGCGGATATGCGCCCTNGATGAAGTCAAGAGGCGATGC 149
.....||| .....|||
48 LysGlnHisIleGlyValAlaGluGluGluLeuGluCysValSerValIleGly 64
150 CGTCAAAAAAGCCCAAGTCTGTT.....GACGCAAAAAGNATCCGG 193
||| .....|||
64 sValLeuArgGlyGlnProLeuThrArgGlyArgGlyLysMetLeuPro. 80
194 GCGTGTGTTTACCGCGCCNGTTTCAGGCAAAATCGCGCCATC..... 237
||| .....|||
81 .....ValHisAlaProThrSerGlyThrValThrAlaIleAlaPro 94
238 .....CATCGCGCGCAAAAAGCGGCTACTGCTCACTGCGTGTGAT 275
||| .....|||
95 HisSerThrAlaHisProSerAlaLeuAlaGluLeu...SerValIleIle 110
276 TGCCTTGAAGCGCAAGAC..... 294
||| .....|||
110 eAspAlaAspGlyGluAspCysTrpIleProArgAspGlyTrpAlaAspT 127
295 .....GAATCGAGTTGCAAGCGTACGCGCCGCAAGCG 327
||| .....|||
127 yTrArgThrArgSerArgGluGluLeuIleGluArgIleHisGlnPheGly 143
||| .....|||
328 TTGGCAACTTAAGCGCGCGCAGNANGNANGCAATCTGATCCATCCGG 377
||| .....|||
144 ValAlaGlyLeuGlyGly.....AlaGlu 151
378 TTGTGTGACTGCGCGGTGATGCGGTTCAGCAAAATCCGCGCGTGG 427
||| .....|||
151 yPheProThrGlyVal.....LysLeuGlnGlyGly 162
428 ATGCGAGCGGTTCGCCATCTGTCATGCGATGACACCAATCCGCTN 477
||| .....|||
162 LysPylsIleGluThrIleIleHisAlaAlaGluGluGluProTyr 178
478 GCGGACAGCCCTGTGTTGATCAAGAAGCGCGCAGCATTTCAAGCG 527
||| .....|||
179 IleThrAlaAspAspArgLeuMetGlnAspCysAlaAlaGlnValAlaGlu 195
528 ANGTNMGCGGTATTTGACCGGTTTGACCGAG..... 558
195 uGlyIleArgIleLeuAlaHisIleLeuGlnProArgGluIleLeuIleG 212
559 .....CGTAAATCCATGTTGTGAAGCAGCGTGGC 588
||| .....|||
212 LylIleGluAspAsnLysProGlnAlaIleSerMetLeuArgAlaValLeu 228
589 GCGACAGCTCGCTGTGAANAATGCTGCCAACATCGAACAACATTCGCG 638
||| .....|||
229 AlaAsp.....SerAsnAspIleSerLeuArgValIlePro 240
639 CGGCGCGCATCGCGCGGT.....TTGAGTG 664
||| .....|||
240 oThrLysTyrProSerGlyValAlaLysGlnLeuThrTyrIleLeuThrG 257
```

665 GCACGACATTCATTTCATTGACCGCGTCGCAACAAACCGTTGG. 714
 257 LysLeuVal.....ProHisGlyArgSerSeraspIle 269
 715 ACCATCAATTCATCAAGATGA.....ATTGCATCGGACGTTTGT 755
 270 GlyValLeuMetClnAsnValGlyThrAlaTyrAlaValLysAlaVala 286
 756 TCACACAGCCGCTGCAACACGACGCGCTGATTCCTTTGGGTGCTC 805
 286 LLeaspGlyGluProIleThrGluArgValValThrLeuThrGlyLys 303
 806 AAGTCAACAAACACGCGCTTCGCTACCGTTTGGGTGCGCAAGTATCG 855
 303 LalleAlaArgProGlyAsnValTyrAlaArgLeuGlyThrProValArg 319
 856 CAATT.....ACTGCGGCGAATTGGTTGACGACGACGCGGAT 899
 320 HisLeuLeuAsnAspAlaGlyPheCysProSerAlaAspGlnMetVal 336
 900 TTCGCGTTGCGTATTGAACGCG 921
 336 eMetGlyGlyProLeuMetGly 343

seq_name: /SIDS1/gcgdata/geneseq/geneseq-emb1/AA2001.DAT:ABG00972

seq_documentation_block:

ID ABG00972 standard; Protein; 4274 AA.
 AC ABG00972;
 DT 13-FEB-2002 (first entry)
 DE Novel human diagnostic protein #963.
 KM Human: chromosome mapping; gene mapping; gene therapy; forensic;
 KM food supplement; medical imaging; diagnostic; genetic disorder.
 OS Homo sapiens.
 PN WO200175067-A2.
 PD 11-OCT-2001.
 PF 30-MAR-2001; 2001WO-US08631.
 PR 31-MAR-2000; 2000US-0540217.
 PR 23-AUG-2000; 2000US-0649167.
 PA (HYSE-) HISEQ INC.
 PI Drmanac RT, Liu C, Tang YN;
 DR N-PSDB; AAS65159.
 PT New isolated polynucleotide and encoded polypeptides, useful in
 PT diagnostics, forensics, gene mapping, identification of mutations
 PT responsible for genetic disorders or other traits and to assess
 PT biodiversity
 XX Claim 20; SEQ ID No 31331; 103pp; English.
 PS
 XX The invention relates to isolated polynucleotide (I) and
 CC polypeptide (II) sequences. (I) is useful as hybridisation probes,
 CC polymerase chain reaction (PCR) primers, oligomers, and for chromosome
 CC and gene mapping, and in recombinant production of (II). The
 CC polynucleotides are also used in diagnostics as expressed sequence tags
 CC for identifying expressed genes. (I) is useful in gene therapy techniques
 CC to restore normal activity of (II) or to treat disease states involving
 CC (II). (II) is useful for generating antibodies against it, detecting or
 CC quantitating a polypeptide in tissue, as molecular weight markers and as
 CC a food supplement. (II) and its binding partners are useful in medical

CC imaging of sites expressing (II). (I) and (II) are useful for treating
 CC disorders involving aberrant protein expression or biological activity.
 CC The polypeptide and polynucleotide sequences have applications in
 CC diagnostics, forensics, gene mapping, identification of mutations
 CC responsible for genetic disorders or other traits to assess biodiversity
 CC and to produce other types of data and products dependent on DNA and
 CC amino acid sequences. ABG00010-ABG30377 represent novel human
 CC diagnostic amino acid sequences of the invention.
 CC Note: The sequence data for this patent did not appear in the printed
 CC specification, but was obtained in electronic format directly from WIPO
 CC at ftp.wipo.int/pub/published_pcr_sequences.
 XX

Sequence 4274 AA;

alignment_scores:
 Quality: 101.50 Length: 455
 Ratio: 0.505 Gaps: 18
 Percent Similarity: 44.176 Percent Identity: 21.538

alignment_block:

US-09-303-518d-127/rev x ABG00972 ..

Align seg 1/1 to: ABG00972 from: 1 to: 4274

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 1495 LysProPhePheSerThrArgProTyrGlnSerThrThrAlaProIle 1511
 1293 TTGCTATTGCGCGGACAGCAAGCTGCACAAAGAGAGCTTCTGCT 1244
 1511 eThr...ValProGlyProAlaLysSerGlyPheThrSerLeuSerSer 1527
 1243 CCAATTCCAAACACCCATGCTGCGCGCTG.....TCGCTA 1206
 1527 eSerSerAsnThrProSerAlaSerProLeuLysSerIleThrSerVal 1543
 1205 TCGCCGACGATTAAATCGCGCAAGGATGAGCAGATGTCAGGCG 1156
 1544 Ser...ThrProSerProIleLysSerThrLeuGlyAlaSerThrThrSe 1559
 1155 CATTACGCGCTCG..... 1143
 1559 rSerValLysSerIleSerAspValAlaSerProIleArgSerLeuArgT 1576
 1143 1143
 1576 hMetSerSerProIleLysThrValValSerGlnSerProTyrAsnIle 1592
 1142 TAAGTACCAATCGACCATGCGCGGTCGCGCCGCTGAGCGGTGCT 1093
 1593 GlnValSerSerGlyThrLeuAlaArgAlaProAlaValThrGlnAlaTh 1609
 1092 GAACCTGAAAGAGTTGTTTTCAGGAATAAGCGGAGGTCTGACCGTGA 1043
 1609 rProLeuLysGlyLeuAlaSerAsnSer..... 1618
 1042 TGGAGATTTCGCGCGCGCGGCGCAACCGCAACGACTTTGCTG 993
 1619 ..ThrPheSerSerArgThrSerProValThrThrAlaGlySerLeuLeu 1634
 992 CGGCTTCCTTCGATACGGAATCTGATTGTGTAGCTCCCAATAATATC 943
 1635 GluArgSerSerIleThr..... 1640
 942 GTGCGCGCCTTGTGTAAATCGCGCGCTTCATATACGACGGAATCAGC 893
 1641MetThrProAlaSerProLysSerAsnIleAsnM 1653
 892 GGTTCCTTCGCTCAACCAATTCGCGCGAGTAATTTGGATACCTTGCA 843
 1653 eTyrSerSerSerLeuProPheLysSerIleIleThrSerAla...Ala 1668

ID	seq_documentation block:	AA
XX	AA059843 standard; Protein; 311	AA.
AC	AA059843;	
XX		
XX		
DT	26-JAN-1995 (first entry)	
XX		
DE	ApoE4Lx2 protease.	
XX		
KW	ApoE4Lx2; protease; enzyme; Alzheimer disease; diagnostic; therapeutic.	
XX		
OS	Homo sapiens.	
XX		
FH	Key	Location/Qualifiers
FT	Domain	
FT	Region	89..104
FT		/label= apoE4 homology
FT	Cleavage-site	35..36
FT	Cleavage-site	43..44
FT	Cleavage-site	182..183
FT	Cleavage-site	295-296
FT	Misc-difference	3
FT		/label= kinase c phosphorylation site
FT	Misc-difference	42
FT		/label= kinase c phosphorylation site
FT	Misc-difference	43
FT		/label= kinase c phosphorylation site
FT	Misc-difference	54
FT		/label= kinase c phosphorylation site
FT	Misc-difference	87
FT		/label= kinase c phosphorylation site
FT	Misc-difference	122
FT		/label= kinase c phosphorylation site
FT	Misc-difference	129
FT		/label= kinase c phosphorylation site
FT	Misc-difference	152
FT		/label= kinase c phosphorylation site
FT	Misc-difference	303
FT		/label= kinase c phosphorylation site
FT	Misc-difference	176..177
FT		/label= casein kinase phosphorylation site
FT	Misc-difference	193..194
FT		/label= casein kinase phosphorylation site
PN	WO9413798-A.	
XX		
PD	23-JUN-1994.	
XX		
PF	16-DEC-1993; 93WO-EP03581.	
XX		
PR	16-DEC-1992; 92CA-2085924.	
PR	04-MAR-1993; 93US-0291401.	
XX		
PA	(BERG/) BERGMANN J E.	
XX		
XX	(PREDD/) PREDDIE R E.	
PI	Bergmann JE, Preddie RE;	
DR	WPI; 1994-234212/28.	
DR	N-PSDB; AA069101.	
XX		
XX	New proteinase esterase-like proteins - used to develop prods.	
PT	for the diagnosis and treatment of Alzheimer's disease and	
PT	related diseases	
PS	Claim 26; Page 44-45; 72pp; English.	
XX		
XX	ApoE4Lx2 is a protease catalyzing the formation of the abnormal beta/A4	
CC	variant of beta-amyloid protein, and is used to develop an inhibitor	
CC	for the diagnosis and treatment of Alzheimer disease, Downs syndrome,	
CC	Parkinson disease, schizophrenia, hyperlipoproteinemia or	
CC	cardiovascular disease.	
XX		

[illegible]


```

801 TTCTCAAGTCACAAACCCAGCCCTTTCGCTACCGTTTGGT... 843
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844 .....GCGAAGTATCGCAATTTACTCG.....GCGGAA 873
294 IuYsAlaIaIaValAlaSerAlaGluSerMetPheAsnAsnGlnGly 310
874 TTGGTTGACCGCAGAACCCGCGATTTCCGGTTGGTATTGAACGGCGC 923
311 ValCysIleAlaProSerArgLeuIle..... 319
924 GATTACACAGCGCGCGATTTATTGGAGCTTACCAATCAGATT 973
320 .ValGluArgSerIleHisLysArgVal.....ValG 330
974 CCGTTATCGAAGAGCCGACGCAAGAGCTTCCGCTGGCTGGCGCG 1023
330 IuIleValAlaIaIaValAlaLysArgArg..... 339
1024 CAGCCGAGCAATATCTCCATCACCCTGACCGCTGGCATTTCTG... 1071
340 GlnProGlyAspProLeuAspProThrThrArgMetGlyAlaLeuValAs 356
1072 .....AAAACAACTCTTCAAGTTC..... 1092
356 PAlaAsnHisAlaAspArgValMetGlyPheIleGlyArgAlaLysAlaA 373
1093 .....ACGACAGCCGTCACAGGTGGCGACCGCGCATGTCGGATT... 1134
373 spGlyAlaThrLeuValAlaGlyGlyThrArgAlaLeuThrGluThrGly 389
1135 GGTACTTACGAGCGCGTAAAGCG..... 1158
390 GlySerTyr.....ValValProThrValPheAspAsnValSerAsnCy 404
1159 .....CTAGCATCTCTGC 1171
404 smetGluIleAlaArgAspGluValPheGlyProValLeuSerValIleP 421
1172 CTACCCCTCTTTGGCGGATTTAATGTCGGCGATACCGACGCGCGCA 1221
421 roValAlaAsnValGlyGluAlaValAlaValAlaAsnAspSerProTyr 437
1222 GCATTGGGTTCTTGGCAATTGGACGACAGAGACCTCG 1258
438 GlyLeuGlyAlaGlyValThrPheAspArgLeuSer 449

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